WEST Search History

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DATE: Thursday, September 09, 2004

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	DB=PGPB,U	SPT,USOC,EPAB,JPAB,DWPI; PLUR=	YES; OP=ADJ
	L19	L15 AND AScr	27
	L18	L17 AND AScr	. 1
	L17	L16 AND PrPSc	95
	L16	L15 AND PrP	581
	L15	prion	4411
	L14	prion	4411
	L13	L12 AND PrPSc	9
	L12	L11 AND PrP	40
	L11	L10 AND prion	273
	L10	514/2,4,12.CCLS.	12236
	L9	L8 AND PrPSc	3
	L8	L7 AND PrP	20
	L7	L6 AND prion	60
	L6	424/185.1.CCLS.	1735
	L5	Schenk.IN.	2901
	L4	Schenk-D-B.IN.	17
	L3	Schenk-D.IN.	7
	L2	Schenk-Dale.IN.	3
	L1	(Schenk-Dale-B.IN.)	43

END OF SEARCH HISTORY

Hit List

Clear Generate Collection Print Fwd Refs Bkwd Refs Generate OACS

Search Results - Record(s) 1 through 27 of 27 returned.

1. Document ID: US 20040147531 A1

Using default format because multiple data bases are involved.

L19: Entry 1 of 27

File: PGPB

Jul 29, 2004

PGPUB-DOCUMENT-NUMBER: 20040147531

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040147531 A1

TITLE: Amidine derivatives for treating amyloidosis

PUBLICATION-DATE: July 29, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Chalifour, Robert J. Ile Bizard CA
Kong, Xianqi Pierrefonds CA
Wu, Xinfu Dollard-des-Ormeaux CA

Lu, Wenshuo LaSalle CA

US-CL-CURRENT: <u>514/256</u>; <u>514/397</u>, <u>514/636</u>

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Affachments | Claims | KMC | Draw Des

2. Document ID: US 20040138178 A1

L19: Entry 2 of 27

File: PGPB

Jul 15, 2004

PGPUB-DOCUMENT-NUMBER: 20040138178

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040138178 A1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

PUBLICATION-DATE: July 15, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Szarek, Walter A. Kingston CA
Kong, Xianqi Pierrefonds CA
Thatcher, Gregory R.J. Kingston CA
Gorine, Boris Edmonton CA

US-CL-CURRENT: <u>514/79</u>; <u>514/114</u>, <u>514/141</u>

ABSTRACT:

h eb bgeeef eb ef be

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

Full Title Citation Front Review	Classification	Date	Reference	Sequences	Attachments	Claims Koon	C Draw Desc

3. Document ID: US 20040048279 A1

L19: Entry 3 of 27

File: PGPB

Mar 11, 2004

PGPUB-DOCUMENT-NUMBER: 20040048279

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040048279 A1

TITLE: Method for detecting methylation states for a toxicological diagnostic

PUBLICATION-DATE: March 11, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47 Olek, Alexander Berlin DE

Piepenbrock, ChristianBerlinDEBerlin, KurtStahnsdorfDE

US-CL-CURRENT: 435/6

ABSTRACT:

The present invention concerns a method for toxicological diagnosis. A DNA sample is taken from an organism or a cell culture, which has previously been subjected to a specific substance that is to be investigated for its toxicological effect. The DNA contained in this sample is chemically pretreated and the base sequence of a part of the modified DNA is determined. A methylation state characteristic for the sample or a characteristic methylation pattern is concluded from this. The effect of a substance on the organism or the cell culture is concluded by comparison with data of the methylation states of other samples and/or compared with other substances from a toxicological point of view.

Full Title Citation Front	Review Classification Date F	Reference Sequences	Attachments Claims	KMC Draw Desc

4. Document ID: US 20040006092 A1

L19: Entry 4 of 27 File: PGPB Jan 8, 2004

PGPUB-DOCUMENT-NUMBER: 20040006092

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20040006092 A1

TITLE: Amidine derivatives for treating amyloidosis

h eb bgeeef eb ef be

Dec 25, 2003

PUBLICATION-DATE: January 8, 2004

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Chalifour, Robert J. Ile Bizard CA
Kong, Xianqi Dollard-des-Ormeaux CA
Wu, Xinfu Dollard-des-Ormeaux CA
Lu, Wenshuo Montreal CA

US-CL-CURRENT: 514/256; 514/397, 514/632

ABSTRACT:

The present invention relates to the use of amidine compounds in the treatment of amyloid-related diseases. In particular, the invention relates to a method of treating or preventing an amyloid-related disease in a subject comprising administering to the subject a therapeutic amount of an amidine compound. Among the compounds for use according to the invention are those according to the following Formula, such that, when administered, amyloid fibril formation, neurodegeneration, or cellular toxicity is reduced or inhibited: 1

Full Title Citation Front Review Classification Date	e Reference Sequences Attachments Claims KMC Draw. Desc	
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File: PGPB

# 5. Document ID: US 20030236392 A1

PGPUB-DOCUMENT-NUMBER: 20030236392

PGPUB-FILING-TYPE: new

L19: Entry 5 of 27

DOCUMENT-IDENTIFIER: US 20030236392 A1

TITLE: Novel full length cDNA

PUBLICATION-DATE: December 25, 2003

# INVENTOR-INFORMATION:

NAME	CITY	STATE	COUNTRY	RULE-47
Isogai, Takao	Ibaraki		JP	
Sugiyama, Tomoyasu	Tokyo		JP	
Otsuki, Tetsuji	Chiba		JP	
Wakamatsu, Ai	Chiba	•	JP	
Sato, Hiroyuki	Osaka		JP	
Ishii, Shizuko	Chiba		JP ·	
Yamamoto, Jun-lchi	Chiba		JP	
lsono, Yuuko	Chiba		JP	
Hio, Yuri	Chiba		JP	
Otsuka, Kaoru	Saitama		JP	
Nagai, Keiichi	Tokyo		JP	
lrie, Ryotaro	Chiba		JP	
Tamechika, lchiro	Osaka		JP	
Seki, Naohiko	Chiba		JP	
Yoshikawa, Tsutomu	Chiba		JP	
Otsuka, Motoyuki	Tokyo		JP	
h eb bgeeef	e b	ef	b e	

Nagahari, Kenji

Tokyo

JP JP

Masuho, Yasuhiko

Tokyo

US-CL-CURRENT: 536/23.1; 435/183, 435/325, 435/6, 435/69.1, 530/350, 702/19

#### ABSTRACT:

Novel full-length cDNAs are provided.

1970 cDNA derived from human have been isolated. The full-length nucleotide sequences of the cDNA and amino acid sequences encoded by the nucleotide sequences have been determined. Because the cDNA of the present invention are full-length and contain the translation start site, they provide information useful for analyzing the functions of the polypeptide.

Full Title Citation Front Review C	lassification Date Reference	Sequences Attachments Cia	ims KNAC Draw Desc
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# 6. Document ID: US 20030232758 A1

L19: Entry 6 of 27

File: PGPB

Dec 18, 2003

PGPUB-DOCUMENT-NUMBER: 20030232758

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030232758 A1

TITLE: Immunological methods and compositions for the treatment of Alzheimer's

disease

PUBLICATION-DATE: December 18, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

St. George-Hyslop, Peter H. Toronto CA
McLaurin, JoAnne Toronto CA

US-CL-CURRENT: 514/12; 435/320.1, 435/325, 435/69.1, 530/324, 536/23.1

#### ABSTRACT:

The present invention relates to immunogenic compositions and peptides comprising residues 4-10 (FRHDSGY) of the amyloid peptide Abeta.sub.42. The invention further relates to antibodies that bind to the Abeta.sub.(4-10) antigenic determinant. The invention provides methods for treating Alzheimer's disease and for reducing the amyloid load in Alzheimers patients. The invention also relates to methods for designing small molecule inhibitors of amyloid deposition.

Full   Title   Citation   Front   Review   Classification   Date   Reference   Sequences   Affachments   Claims   KMC   Draw Desc

# 7. Document ID: US 20030185808 A1

L19: Entry 7 of 27

File: PGPB

Oct 2, 2003

PGPUB-DOCUMENT-NUMBER: 20030185808

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e b

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PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030185808 A1

TITLE: Prostate cell lines

PUBLICATION-DATE: October 2, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Thraves, Peter London GB Sutton, Andrew London GB

US-CL-CURRENT: 424/93.21; 424/85.2, 435/366, 514/44

#### ABSTRACT:

An increasingly aged population and better diagnosis has lead to an apparent increase in the prevalence of prostate cancer in men. There is an acute need to better understand the progression of this disease from its locally confined site of initiation to the end stage widely metastatic disease with attendant morbidity and mortality. It has historically been difficult to raise and maintain immortalised prostate cell lines in culture. We have derived a cell line selected from the group consisting of clones ONYCAP 1 and ONYCAP23. The cell lines are characterised as being prostate epithelial in origin.

Full Title Citation Front Review	Classification Date	Reference Sequences	Atlachments Claims	KWMC - Draw, Desc
				-

8. Document ID: US 20030108595 A1

L19: Entry 8 of 27 File: PGPB Jun 12, 2003

PGPUB-DOCUMENT-NUMBER: 20030108595

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030108595 A1

TITLE: Method for treating amyloidosis

PUBLICATION-DATE: June 12, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Kisilevsky, Robert Kingston CA Szarek, Walter Kingston CA Weaver, Donald Kingston CA

US-CL-CURRENT:  $\underline{424/450}$ ;  $\underline{514/12}$ ,  $\underline{514/23}$ ,  $\underline{514/378}$ ,  $\underline{514/381}$ ,  $\underline{514/460}$ ,  $\underline{514/79}$ 

# ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates

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and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

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Full Title Citation Front Review Classification Date Reference Sequences Attachments Claims KWIC Draw Des-

9. Document ID: US 20030027796 A1

L19: Entry 9 of 27

File: PGPB

Feb 6, 2003

PGPUB-DOCUMENT-NUMBER: 20030027796

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20030027796 A1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

PUBLICATION-DATE: February 6, 2003

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Szarek, Walter A. Kingston CA

Kong, Xianqi Dollard-des-Ormeaux CA
Thatcher, Gregory R.J. Kingston CA

Gorine, Boris Edmonton CA

US-CL-CURRENT: 514/79; 514/114, 514/141

# ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

Full Title Citation Front Review Classificat	ion Date Reference Sequ	uences   Attachments   Claims	KMMC Draw Desc
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10. Document ID: US 20020119926 A1

L19: Entry 10 of 27 File: PGPB Aug 29, 2002

PGPUB-DOCUMENT-NUMBER: 20020119926

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020119926 A1

TITLE: Inhibitors of IAPP fibril formation and uses thereof

PUBLICATION-DATE: August 29, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

h eb bgeeef eb ef be

Jul 18, 2002

Fraser, Paul

Toronto

CA

US-CL-CURRENT: <u>514/12</u>; <u>435/184</u>, <u>514/14</u>, <u>514/15</u>, <u>514/16</u>, <u>514/17</u>

#### ABSTRACT:

New antifibrillogenic agents and compositions containing same, methods of using the antifibrillogenic agents and compositions for inhibiting amyloid fibril formation, and effective therapeutics for preventing or delaying the progression of, e g., Alzheimer's disease and diabetes.

. Full∷	Title Citation	Front Review	Classification   Date	a Reference	Sequences	Attachments	Claims	KMC	Draw Desc
	11. Docur	ment ID: US 2	0020115717 A	1	***************************************		***************************************		
L19:	Entry 11 o	f 27		File:	PGPB		Aug	22,	2002

PGPUB-DOCUMENT-NUMBER: 20020115717

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020115717 A1

TITLE: Amyloid targeting imaging agents and uses thereof

PUBLICATION-DATE: August 22, 2002

#### INVENTOR-INFORMATION:

CITY	STATE	COUNTRY	RULE-47
Ile Bizard		CA	
Dollard-des-Ormeaux		CA	
Ile Bizard		CA	
Laval		CA	
	Ile Bizard Dollard-des-Ormeaux Ile Bizard	Ile Bizard Dollard-des-Ormeaux Ile Bizard	Ile Bizard CA Dollard-des-Ormeaux CA Ile Bizard CA

US-CL-CURRENT: <u>514/553</u>; <u>424/1.11</u>

#### ABSTRACT:

Amyloid-targeting imaging agents such as radiolabeled amyloid targeting molecules and amyloid targeting molecule-chelator conjugates for imaging, e.g., amyloid plaques in vivo, and/or for the treatment of amyloidosis disorders. The invention provides amyloid-targeting imaging agents that are useful for imaging sites of amyloid disease. Imaging agents of the invention are capable of binding specifically to amyloid plaques, as an aid in diagnosis and/or early treatment of amyloidosis disorders.

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Fulf	Title	Citation Front	Review	Classification	Date	Reference	Sequences	Attachments	Claims	KOMC	Drawn Desc
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1000	14.	Document ID	. USZ	002009433	JAI						

File: PGPB

PGPUB-DOCUMENT-NUMBER: 20020094335

PGPUB-FILING-TYPE: new

L19: Entry 12 of 27

DOCUMENT-IDENTIFIER: US 20020094335 A1

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TITLE: Vaccine for the prevention and treatment of alzheimer's and amyloid related diseases

PUBLICATION-DATE: July 18, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Chalifour, Robert Ile Bizard CA
Hebert, Lise Brossard CA

Kong, Xianqi Dollard-des-Oremaux CA Gervais, Francine Ile Bizard CA

US-CL-CURRENT: <u>424/185.1</u>

# **ABSTRACT:**

The present invention relates to a stereochemically based "non-self" antigen vaccine for the prevention and/or treatment of Alzheimer's and other amyloid related diseases. The present invention provides a vaccine for the prevention and treatment of Alzheimer's and other amyloid related diseases, which overcomes the drawbacks associated with using naturally occurring peptides, proteins or immunogens.

Full Title Citation Front Review	Classification Date	Reference	Sequences	Attachments	Claims KM0	Draw Desc

13. Document ID: US 20020009730 A1

L19: Entry 13 of 27 File: PGPB Jan 24, 2002

PGPUB-DOCUMENT-NUMBER: 20020009730

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20020009730 A1

TITLE: Human stress array

PUBLICATION-DATE: January 24, 2002

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Chenchik, Alex Palo Alto CA US

Lukashev, Matvey E. Newton MA US

US-CL-CURRENT: 435/6; 536/24.3

## ABSTRACT:

Human stress arrays and methods for their use are provided. The subject arrays include a plurality of polynucleotide spots, each of which is made up of a polynucleotide probe composition of unique polynucleotides corresponding to a human stress gene. The subject arrays find use in hybridization assays, particularly in assays for the identification of differential gene expression of human stress genes.

Full Title Citation Front	Review Classification Date	Reference Sequences	Attachments   Claims   KWC   Draw	Desc
				·······

14. Document ID: US 20010048941 A1

L19: Entry 14 of 27

File: PGPB

Dec 6, 2001

PGPUB-DOCUMENT-NUMBER: 20010048941

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20010048941 A1

TITLE: Method for treating amyloidosis

PUBLICATION-DATE: December 6, 2001

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Kisilevsky, Robert Kingston CA Szarek, Walter Kingston CA Weaver, Donald Kingston CA

US-CL-CURRENT: 424/450; 514/2, 514/378, 514/381, 514/460, 514/54

## ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amnyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof

Full Title Citation Front Review	Classification Date Reference	e Sequences Attachments Clair	ns   KWMC   Draw Desc

## 15. Document ID: US 20010027186 A1

L19: Entry 15 of 27

File: PGPB

Oct 4, 2001

PGPUB-DOCUMENT-NUMBER: 20010027186

PGPUB-FILING-TYPE: new

DOCUMENT-IDENTIFIER: US 20010027186 A1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

PUBLICATION-DATE: October 4, 2001

INVENTOR-INFORMATION:

NAME CITY STATE COUNTRY RULE-47

Szarek, Walter A. Kingston CA
Kong, Xianqi Dollard-des-Ormeaux CA
Thatcher, Gregory R.J Kingston CA
Gorine, Boris Edmonton CA

US-CL-CURRENT: 514/79; 514/114, 514/129, 514/142

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## ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

Full | Title | Citation | Front | Review | Classification | Date | Reference | Sequences | Attachments | Claims | KNAC | Draw, Desc

16. Document ID: US 6632808 B1

L19: Entry 16 of 27

File: USPT

Oct 14, 2003

US-PAT-NO: 6632808

DOCUMENT-IDENTIFIER: US 6632808 B1

TITLE: Inhibitors of amyloid formation

DATE-ISSUED: October 14, 2003

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Caughey; Winslow S.

Hamilton

MT

Caughey; Byron

Hamilton

МТ

US-CL-CURRENT: 514/185; 514/410, 540/122, 540/145

#### ABSTRACT:

Methods, compounds and compositions are disclosed for treating amyloidogenic diseases, like Alzheimer's disease and type 2 diabetes, and particularly <u>prion</u> diseases associated with conversion of protease sensitive PrP (PrP-sen) to protease resistant PrP (PrP-res), by administering therapeutically effective amounts of a tetrapyrrole. Particular disclosed tetrapyrroles having this activity include phthalocyanines, deuteroporphyrins, and meso-substituted porphines. Complexes of certain of the pyrroles with metals or metal ions produce compounds that are particularly effective in converting the conversion of PrP-sen to PrP-sen. The treatment of the present invention is particularly suited for preventing or inhibiting the progression of <u>prion</u> related diseases, such as transmissible spongiform encephalopathies.

70 Claims, 10 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 4

Full Title: Citation Front Review Classification Date Reference

17. Document ID: US 6562836 B1

L19: Entry 17 of 27

File: USPT

May 13, 2003

US-PAT-NO: 6562836

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DOCUMENT-IDENTIFIER: US 6562836 B1

** See image for Certificate of Correction **

TITLE: Methods and compounds for inhibiting amyloid deposits

DATE-ISSUED: May 13, 2003

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY

Szarek; Walter A. Kingston CA Weaver; Donald F. Kingston CA Kong; Xianqi Dollard-des-Ormeaux CA Gupta; Ajay Pointe-Claire CA Migneault; David Laval CA

US-CL-CURRENT: 514/307; 514/308, 514/311, 514/313, 514/314

#### ABSTRACT:

Methods and compositions which are useful in the treatment of amyloidosis. In particular, methods and compositions are provided for inhibiting, preventing and treating amyloid deposition, e.g., in pancreatic islets, wherein the amyloidotic deposits are islet amyloid polypeptide (IAPP)-associated amyloid deposition or deposits. The methods of the invention involve administering to a subject a therapeutic compound which inhibits IAPP-associated amyloid deposits. Accordingly, the compositions and methods of the invention are useful for inhibiting IAPPassociated amyloidosis in disorders in which such amyloid deposition occurs, such as diabetes.

172 Claims, 14 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 14

Full Titl	e   Citation   Front   Rev	ew   Classification   Date	Reference	Claims  KMC   I	Drawi Desi
□ 18	Document ID: U	S 6440952 B2		 	,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,

L19: Entry 18 of 27

File: USPT Aug 27, 2002

US-PAT-NO: 6440952

DOCUMENT-IDENTIFIER: US 6440952 B2

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

DATE-ISSUED: August 27, 2002

INVENTOR-INFORMATION:

CITY STATE ZIP CODE COUNTRY NAME Szarek; Walter A. Kingston CA CA Kong; Xianqi Dollard-des-Ormeaux Thatcher; Gregory R. J. Kingston CA Gorine; Boris Edmonton CA

US-CL-CURRENT: 514/120; 558/110, 558/70

h e b b g ee e f ef ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

20 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full Title Citation Front Review Classifica	tion Date Reference	Claims   WMC   Draw Desi
19. Document ID: US 6355784	B1	
L19: Entry 19 of 27	File: USPT	Mar 12, 2002

US-PAT-NO: 6355784

DOCUMENT-IDENTIFIER: US 6355784 B1

** See image for <u>Certificate of Correction</u> **

TITLE: Methods and compositions for the manufacture of halogenated anthracyclines with increased antitumor activity, other anthracyclines, halogenated sugars, and glycosyl donors

DATE-ISSUED: March 12, 2002

# INVENTOR-INFORMATION:

NAME	CITY	STATE	ZIP CODE	COUNTRY
Priebe; Waldemar	Houston	TX	77005	
Krawczyk; Marta	Lexington	KY	40503	
Skibicki; Piotr	Warsaw 04015			PL
Fokt; Izabela	The Woodlands	TX	77380	
Dziewiszek; Krzysztof	The Woodlands	TX	77380	
Grynkiewicz; Grzegorz	05-092 Lomianki			PL
Perez-Soler; Roman	New York	NY	10016	

US-CL-CURRENT:  $\underline{536}/\underline{6.4}$ ;  $\underline{536}/\underline{122}$ ,  $\underline{536}/\underline{17.2}$ ,  $\underline{536}/\underline{18.4}$ ,  $\underline{536}/\underline{18.7}$ ,  $\underline{536}/\underline{4.1}$ 

#### ABSTRACT:

The present invention discloses new and novel halogenated anthracyclines linked through the saccharide portions. These congeners show high activity in vitro against several tumor cell lines. In doxorubicin (DOX) sensitive cell lines, they are at least as cytotoxic as DOX and in some cases more so. Many of these 4'- and 6'-fluorinated anthracyclines are more effective against multidrug-resistant tumors than was DOX, and/or have greater effectiveness than DOX against DOX sensitive cells. The compounds of this invention also have anti-amyloidogenic effects and the use of these compounds in the treatment of Alzheimer's disease is contemplated.

7 Claims, 19 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 15 Full : Title: Citation Front: Review Classification Date Reference

20. Document ID: US 6329356 B1

L19: Entry 20 of 27

File: USPT

Dec 11, 2001

US-PAT-NO: 6329356

DOCUMENT-IDENTIFIER: US 6329356 B1

TITLE: Phosphono-carboxylate compounds for treating amyloidosis

DATE-ISSUED: December 11, 2001

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Szarek; Walter A.

Kong; Xiangi

Kingston
Dollard-des-Ormeaux

CA

CA

US-CL-CURRENT: <u>514/120</u>

### ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, a congener thereof, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

31 Claims, 0 Drawing figures Exemplary Claim Number: 1

Full Title Citation Front Review Classification Date Reference	Claims   KWC   Draw Desc

# 21. Document ID: US 5972328 A

L19: Entry 21 of 27

File: USPT

Oct 26, 1999

US-PAT-NO: 5972328

DOCUMENT-IDENTIFIER: US 5972328 A

** See image for <u>Certificate of Correction</u> **

TITLE: Method for treating amyloidosis

DATE-ISSUED: October 26, 1999

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Kisilevsky; Robert

Kingston

CA

Szarek; Walter

Kingston

CA

Weaver; Donald

Kingston

CA

US-CL-CURRENT:  $\underline{424/78.31}$ ;  $\underline{424/423}$ ,  $\underline{424/427}$ ,  $\underline{424/430}$ ,  $\underline{424/434}$ ,  $\underline{424/436}$ ,  $\underline{424/441}$ ,

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<u>424/450</u>, <u>424/78.35</u>, <u>514/772.4</u>, <u>526/286</u>, <u>526/287</u>

#### ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

58 Claims, 10 Drawing figures Exemplary Claim Number: 1
Number of Drawing Sheets: 8

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# 22. Document ID: US 5869469 A

L19: Entry 22 of 27

File: USPT

Feb 9, 1999

US-PAT-NO: 5869469

DOCUMENT-IDENTIFIER: US 5869469 A

TITLE: Phosphonocarboxylate compounds for treating amyloidosis

DATE-ISSUED: February 9, 1999

INVENTOR-INFORMATION:

NAME

CITY

STATE ZIP CODE

COUNTRY

Szarek; Walter A.

Kingston

CA

Kong; Xianqi

Kingston

 ca 

US-CL-CURRENT: 514/120

#### ABSTRACT:

Therapeutic compounds and methods for modulating amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is modulated by the administration to a subject of an effective amount of a therapeutic compound comprising a phosphonate group and a carboxylate group, or a pharmaceutically acceptable salt or ester thereof. In preferred embodiments, an interaction between an amyloidogenic protein and a basement membrane constituent is modulated.

25 Claims, 0 Drawing figures Exemplary Claim Number: 1

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23. Document ID: US 5858326 A

L19: Entry 23 of 27

File: USPT

Jan 12, 1999

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Nov 24, 1998

# Record List Display

US-PAT-NO: 5858326

DOCUMENT-IDENTIFIER: US 5858326 A

TITLE: Methods of increasing amyloid deposition

DATE-ISSUED: January 12, 1999

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Kisilevsky; Robert Kingston CA Szarek; Walter Kingston CA Weaver; Donald Kingston CA Fraser; Paul Toronto CA Kong; Xianqi Kingston CA

US-CL-CURRENT:  $\underline{424/9.2}$ ;  $\underline{424/78.31}$ ,  $\underline{424/78.35}$ ,  $\underline{435/7.8}$ ,  $\underline{435/7.92}$ ,  $\underline{435/7.93}$ ,  $\underline{435/7.95}$ , <u>514/772.4</u>, <u>530/350</u>, <u>800/9</u>

#### ABSTRACT:

In vivo and in vitro methods of increasing amyloid deposition using amyloid-enhancing compounds are described. Methods of forming amyloid fibrils and screening for agents useful in treating amyloidosis are also described. Animals having non-naturally occurring amyloid deposits produced using the amyloid-enhancing compounds even further are described.

5 Claims, 2 Drawing figures Exemplary Claim Number: 5 Number of Drawing Sheets: 2

Full	Title	Citation   Front	Review Classification	Date Reference		Claims	CMC -	Drawn Desi
	24	Document ID	: US 5840294 A		,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,,	***************************************	)	***************************************
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US-PAT-NO: 5840294

DOCUMENT-IDENTIFIER: US 5840294 A

** See image for <u>Certificate of Correction</u> **

TITLE: Method for treating amyloidosis

DATE-ISSUED: November 24, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY Kisilevsky; Robert Kingston CA Szarek; Walter Kingston CA Weaver: Donald Kingston CA

US-CL-CURRENT: 424/78.31; 424/423, 424/427, 424/430, 424/434, 424/436, 424/441, <u>424/450, 424/78.35, 514/772.4, 526/286, 526/287</u>

ABSTRACT:

h b g ee e f e b e b ef Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

66 Claims, 14 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 12

Full	Title Citation Front Review Classification Date	Reference CI.	aims KWMC Drawn Desc
	25. Document ID: US 5728375 A	······	**************************************
•	Entry 25 of 27	File: USPT	Mar 17, 1998

US-PAT-NO: 5728375

DOCUMENT-IDENTIFIER: US 5728375 A

TITLE: Method for treating amyloidosis

DATE-ISSUED: March 17, 1998

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY
Kisilevsky; Robert Kingston CA
Szarek; Walter Kingston CA
Weaver; Donald Kingston CA

US-CL-CURRENT: 424/78.31; 424/450, 424/78.35, 514/772.4, 526/286, 526/287

# ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

71 Claims, 12 Drawing figures Exemplary Claim Number: 1
Number of Drawing Sheets: 12

Full Title	Citation Front Review Classification Date Reference
26.	Document ID: US 5643562 A

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L19: Entry 26 of 27

File: USPT

Jul 1, 1997

US-PAT-NO: 5643562

DOCUMENT-IDENTIFIER: US 5643562 A

TITLE: Method for treating amyloidosis

DATE-ISSUED: July 1, 1997

INVENTOR-INFORMATION:

NAME CITY STATE ZIP CODE COUNTRY
Kisilevsky; Robert Kingston CA
Szarek; Walter Kingston CA
Weaver; Donald Kingston CA

US-CL-CURRENT: 424/78.31; 424/423, 424/427, 424/430, 424/434, 424/436, 424/441, 424/78.35, 514/772.4, 526/286, 526/287

#### ABSTRACT:

Therapeutic compounds and methods for inhibiting amyloid deposition in a subject, whatever its clinical setting, are described. Amyloid deposition is inhibited by the administration to a subject of an effective amount of a therapeutic compound comprising an anionic group and a carrier molecule, or a pharmaceutically acceptable salt thereof, such that an interaction between an amyloidogenic protein and a basement membrane constituent is inhibited. Preferred anionic groups are sulfonates and sulfates. Preferred carrier molecules include carbohydrates, polymers, peptides, peptide derivatives, aliphatic groups, alicyclic groups, heterocyclic groups, aromatic groups and combinations thereof.

55 Claims, 12 Drawing figures Exemplary Claim Number: 1
Number of Drawing Sheets: 12

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27. Document ID: US 5276059 A

L19: Entry 27 of 27

File: USPT

Jan 4, 1994

US-PAT-NO: 5276059

DOCUMENT-IDENTIFIER: US 5276059 A

** See image for <u>Certificate of Correction</u> **

TITLE: Inhibition of diseases associated with amyloid formation

DATE-ISSUED: January 4, 1994

INVENTOR-INFORMATION:

NAME CITY

STATE

ZIP CODE

COUNTRY

Caughey; Byron

Hamilton

MT

Race; Richard

Hamilton

MT

US-CL-CURRENT: 514/647

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ABSTRACT:

The invention provides a method of treating a mammal having a condition associated with formation of amyloidogenic protein without deposition of amyloid plaques. This treatment includes administering to the mammal a pharmacologically effective amount of Congo Red or a pharmaceutically acceptable salt or derivative thereof to interfere with amyloidogenic protein formation or to destabilize amyloidogenic protein structures already formed in said mammal. The invention also provides a method of treating a mammal having a condition associated with deposition of amyloidogenic protein in plaques, and a method of inhibiting the transformation of PrP-sen to PrP-res in a tissue culture sample containing PrP-sen.

34 Claims, 4 Drawing figures Exemplary Claim Number: 1 Number of Drawing Sheets: 3

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J Infect Dis. 2004 Feb 1;189(3):431-9. Epub 2004 Jan 21. PMID: 14745700 [PubMed - indexed for MEDLINE]

2: Zanusso G, Casalone C, Acutis P, Bozzetta E, Farinazzo A, Gelati M, Fiorini M, Forloni G, Sy MS, Monaco S, Caramelli M.

Molecular analysis of iatrogenic scrapie in Italy.
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PMID: 12655108 [PubMed - indexed for MEDLINE]

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Synthetic peptide vaccines yield monoclonal antibodies to cellular and pathological prion proteins of ruminants.

J Gen Virol. 1998 Apr;79 ( Pt 4):937-45. PMID: 9568991 [PubMed - indexed for MEDLINE]

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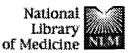
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Immunization delays the onset of prion disease in mice.

Sigurdsson EM, Brown DR, Daniels M, Kascsak RJ, Kascsak R, Carp R, Meeker HC, Frangione B, Wisniewski T.

Department of Psychiatry, New York University School of Medicine, New York, New York 10016, USA.

The outbreak of new variant Creutzfeldt-Jakob disease has raised the specter of a potentially large population being at risk to develop this prionosis. None of the prionoses currently have an effective treatment. Recently, vaccination has been shown to be effective in mouse models of another neurodegenerative condition, namely Alzheimer's disease. Here we report that vaccination with recombinant mouse prion protein delays the onset of prion disease in mice. Vaccination was performed both before peripheral prion exposure and after exposure. A delay in disease onset was seen in both groups, but was more prolonged in animals immunized before exposure. The increase in the incubation period closely correlated with the anti-prion protein antibody titer. This promising finding suggests that a similar approach may work in humans or other mammalian species at risk for prion disease.

PMID: 12107084 [PubMed - indexed for MEDLINE]

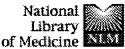
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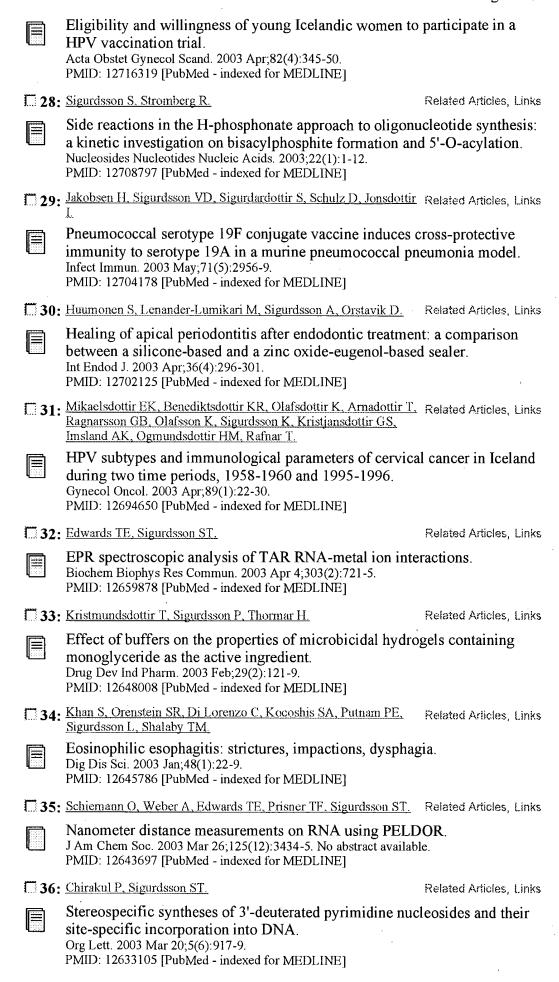
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Magn Reson Med. 2003 Aug;50(2):293-302. PMID: 12876705 [PubMed - indexed for MEDLINE] 19: Sigurdsson S. Richer HB. Hansen BM, Stairs IH. Thorsett SE. Related Articles, Links A young white dwarf companion to pulsar B1620-26: evidence for early planet formation. Science. 2003 Jul 11;301(5630):193-6. PMID: 12855802 [PubMed] 20: Revnisdottir I, Thorleifsson G, Benediktsson R, Sigurdsson G, Related Articles, Links Emilsson V, Einarsdottir AS, Hjorleifsdottir EE, Orlygsdottir GT, Bjornsdottir GT, Saemundsdottir J, Halldorsson S, Hrafnkelsdottir S. Sigurjonsdottir SB, Steinsdottir S, Martin M, Kochan JP, Rhees BK, Grant SF, Frigge ML, Kong A, Gudnason V, Stefansson K, Gulcher JR. Localization of a susceptibility gene for type 2 diabetes to chromosome 5q34-q35.2. Am J Hum Genet. 2003 Aug;73(2):323-35. Epub 2003 Jul 08. PMID: 12851856 [PubMed - indexed for MEDLINE] 21: Dovere V. Schafe GE, Sigurdsson T, LeDoux JE. Related Articles, Links Long-term potentiation in freely moving rats reveals asymmetries in thalamic and cortical inputs to the lateral amygdala. Eur J Neurosci. 2003 Jun;17(12):2703-15. PMID: 12823477 [PubMed - indexed for MEDLINE] 22: Sigurdardottir M, Sigurdsson H, Barkardottir RB, Kristjansdottir S, Related Articles, Links Agnarsson BA. Lymphoid tumours of the ocular adnexa: a morphologic and genotypic = study of 15 cases. Acta Ophthalmol Scand. 2003 Jun;81(3):299-303. PMID: 12780412 [PubMed - indexed for MEDLINE] 23: Sigurdsson G, Yannopoulos D, McKnite SH, Lurie KG. Related Articles, Links Cardiorespiratory interactions and blood flow generation during cardiac arrest and other states of low blood flow. Curr Opin Crit Care. 2003 Jun;9(3):183-8. Review. PMID: 12771667 [PubMed - indexed for MEDLINE] 24: van Kooij B, van Dijk MC, de Boer J, Sigurdsson V, Rothova A. Related Articles, Links Is granuloma annulare related to intermediate uveitis with retinal vasculitis? Br J Ophthalmol. 2003 Jun;87(6):763-6. PMID: 12770977 [PubMed - indexed for MEDLINE] 25: Sigurdsson U, Adolphson P. Related Articles, Links [Few injuries among Swedish telemark skiers, but equipment requires careful consideration] Lakartidningen. 2003 Apr 17;100(16):1440-3. Swedish. PMID: 12756712 [PubMed - indexed for MEDLINE] 26: Borda EJ. Markley JC. Sigurdsson ST. Related Articles, Links Zinc-dependent cleavage in the catalytic core of the hammerhead ribozyme: evidence for a pH-dependent conformational change. Nucleic Acids Res. 2003 May 15;31(10):2595-600. PMID: 12736309 [PubMed - indexed for MEDLINE] 7: Gudmundsdottir T, Tryggvadottir L, Allende M, Mast TC, Briem Related Articles, Links H. Sigurdsson K.



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□ 38	Sigfusson N, Sigurdsson G, Agnarsson U, Gudmundsdottir II, Stefansdottir I, Sigvaldason H, Gudnason V.	Related Articles, Links
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	Functional cross-talk among Rad51, Rad54, and replical heteroduplex DNA joint formation.  J Biol Chem. 2002 Nov 15;277(46):43578-87. Epub 2002 Sep 10.  PMID: 12226081 [PubMed - indexed for MEDLINE]	tion protein A in
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	Infectivity of amyloid diseases. Trends Mol Med. 2002 Sep;8(9):411-3. PMID: 12223307 [PubMed - indexed for MEDLINE]	
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Anti-prion antibodies for prophylaxis following prion exposure in mice.

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Sigurdsson EM, Sy MS, Li R, Scholtzova H, Kascsak RJ, Kascsak R, Carp R, Meeker HC, Frangione B, Wisniewski T.

Department of Psychiatry, New York University School of Medicine, Millhauser Laboratory, 550 First Avenue, New York, NY 10016, USA.

Prion disease is characterized by a conformational change of the normal form of the prion protein (PrP(C)) to the scrapie-associated form (PrP(Sc)). Since the emergence of new variant Creutzfeldt-Jakob disease a potentially large human population is at risk for developing prion disease. Currently, no effective treatment or form of post-exposure prophylaxis is available for prion disease. We recently showed that active immunization with recombinant PrP prolongs the incubation period of scrapie. Here we show that anti-PrP antibodies following prion exposure are effective at increasing the incubation period of the infection. Stimulation of the immune system is an important therapeutic target for the prion diseases, as well as for other neurodegenerative illnesses characterized by abnormal protein conformation.

PMID: 12505623 [PubMed - indexed for MEDLINE]

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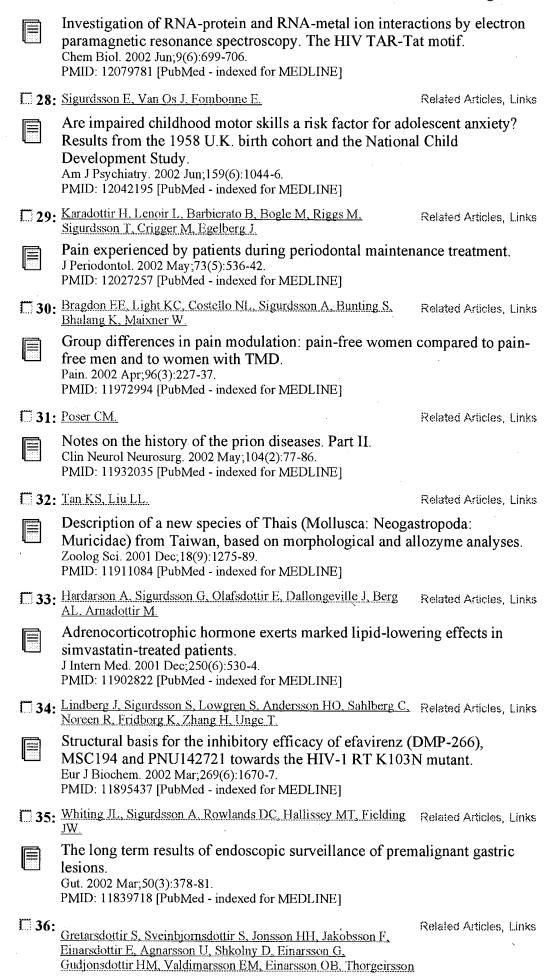
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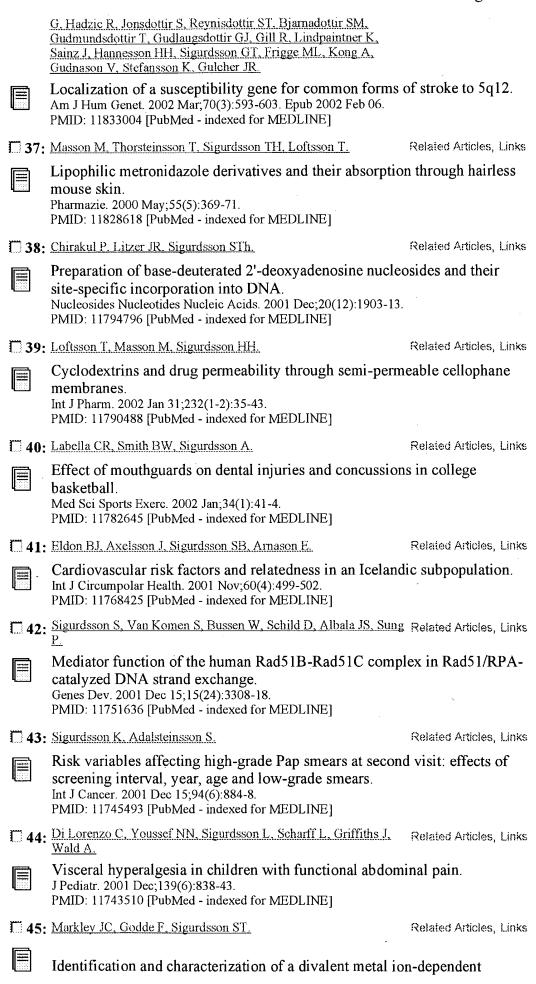


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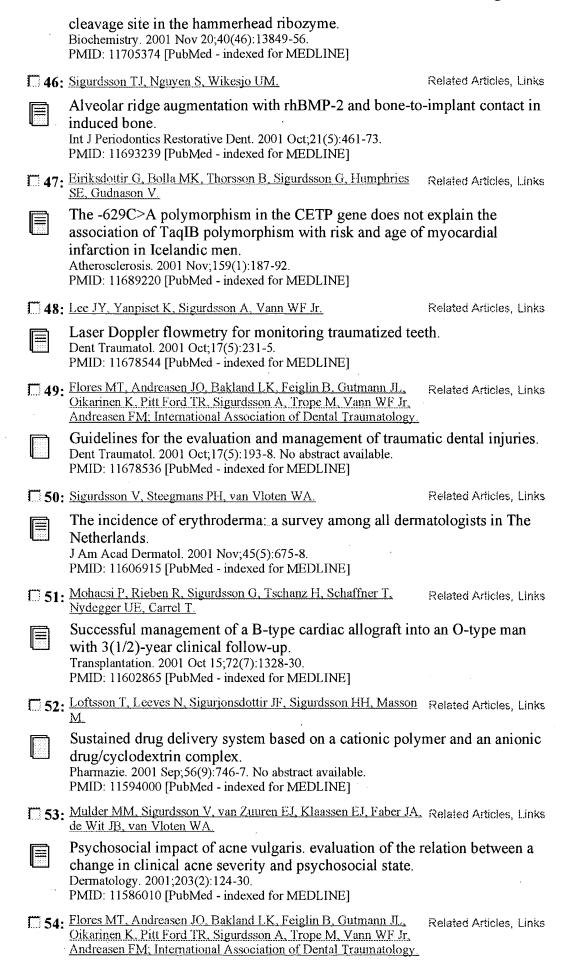
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Synthetic peptide vaccines yield monoclonal antibodies to cellular and pathological prion proteins of ruminants.

Harmeyer S, Pfaff E, Groschup MH.

Federal Research Centre for Virus Diseases of Animals, Tubingen, Germany.

Transmissible spongiform encephalopathies are closely linked to the accumulation of a pathological isoform of a host-encoded prion protein (PrP (C)), designated PrP(Sc). In an attempt to generate mono- and polyclonal antibodies to ruminant PrP, 32 mice were vaccinated with peptide vaccines which were synthesized according to the amino acid sequence of ovine PrP. By this approach five PrP-reactive polyclonal antisera directed against four different domains of the protein were stimulated. Splenocytes of mice which had developed PrP-reactive antibodies were used for the generation of monoclonal antibodies (MAbs). Obtained PrP-specific MAbs were directed to three different domains of ruminant PrP which differed from the three previously described major MAb binding sites in rodent PrP. MAbs exhibited reactivity with non-denatured ruminant PrP(C) in ELISA and immunoprecipitation and with denatured ovine and bovine PrP(Sc) in immunoblot. Cross-reactivity was observed with PrP(C) of nine other mammalian species and with pathological PrP preferably of ruminants and weakly with that of hamster and mouse. The generated MAbs will be useful tools for the development of diagnostic tests for BSE and scrapie as well as for pathogenesis studies of these diseases.

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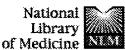
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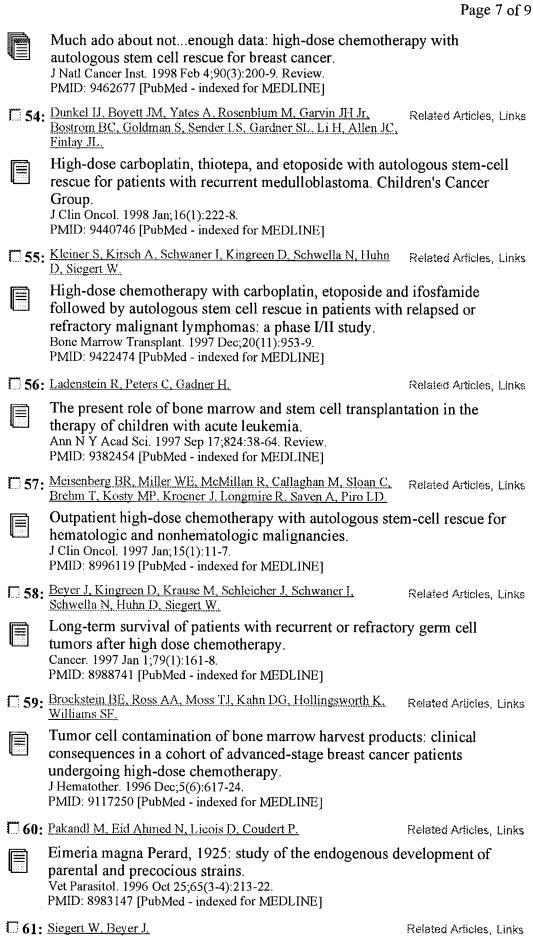
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      us 2002-169492
                             A2
                                    20021021
L3
      ANSWER 3 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
      2004:141667
ΑN
                    CAPLUS
      140:198062
DN
TI
      Antigen arrays and compositions comprising antigenic determinant of RANKL
      protein and virus-like particle for treatment of bone disease and prion
IN
      Bachmann, Martin; Maurer, Patrick; Spohn, Gunther
PA
      U.S. Pat. Appl. Publ., 53 pp., Cont.-in-part of Appl. No. PCT/IB02/00166.
S0
      CODEN: USXXCO
      Patent
      English
FAN.CNT 9
      PATENT NO.
                            KIND
                                    DATE
                                                 APPLICATION NO.
                                                                           DATE
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                                                 ______
PΙ
      US 2004033211
                                    20040219
                                                 US 2002-289456
                            Α1
                                                                           20021107
                                                 US 2002-50902
      US 2003175290
                             Α1
                                    20030918
                                                                           20020118
     WO 2002056905
                             A2
                                    20020725
                                                 WO 2002-IB166
                                                                           20020121
     wo 2002056905
                             Α3
                                    20031009
              AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
               CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
              PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM,
                                                                    TN, TR,
                                                                             TT, TZ,
              UA, UG, US, UZ, VN, YU, ZA, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU,
          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH,
              CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR,
               BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG
PRAI US 2001-331045P
                            Р
                                    20011107
      us 2002-50902
                             A2
                                    20020118
     WO
        2002-IB166
                             Α2
                                    20020121
        2002-396635P
     US
                             Ρ
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     US 2001-262379P
                             Ρ
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     US 2001-288549P
                             Ρ
                                    20010504
     US 2001-326998P
                                    20011005
L3
     ANSWER 4 OF 76 USPATFULL ON STN
        2004:171948 USPATFULL
AN
ΤI
        Method
IN
        Enari, Masato, Chuo-ku, JAPAN
        Flechsig, Eckhard, Versbacher, GERMANY, FEDERAL REPUBLIC OF Collinge, John, Queen, UNITED KINGDOM
       Weismann, Charles, London, UNITED KINGDOM
       US 2004132109
                                   20040708
PΙ
                             Α1
        us 2004-470022
                                   20040109 (10)
ΑI
                             Α1
       WO 2002-GB257
                                   20020122
PRAI
        GB 2001-1762
                              20010123
DT
       Utility
FS
        APPLICATION
LN.CNT 3141
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INCL
          INCLM: 435/007.200
          INCLS: 435/287.200
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          NCLM:
                   435/007.200
          NCLS:
                  435/287.200
          [7]
 IC
          ICM: G01N033-53
          ICS: G01N033-567; C12M001-34
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
 L3
       ANSWER 5 OF 76 USPATFULL on STN
          2004:94784 USPATFULL
AN
            ***PrPSc***
                               -interacting molecules and uses thereof
ΤI
          Cashman, Neil, Toronto, CANADA
IN
          Paramithiotis, Eustache, Boucherville, CANADA
          La Boissiere, Sylvie, Montreal, CANADA
         Lawton, Robert, Gorham, ME, UNITED STATES
Francoeur, Greg, North Yormouth, ME, UNITED STATES
         Francoeur, Susan, Portland, ME, UNITED STATES
          Estey, Lisa, Westbrook, ME, UNITED STATES
          Pinard, Marc, Montreal, CANADA
         US 2004072236
PΙ
                                  Α1
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ΑI
         us 2002-256538
                                  Α1
                                         20020927 (10)
         Utility
DT
         APPLICATION
FS
LN.CNT 1436
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         NCLM:
                 435/007.100
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IC
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CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
       ANSWER 6 OF 76 USPATFULL on STN
         2004:69593 USPATFULL
ΑN
         Fusion proteins comprising DP-178 and other viral fusion inhibitor peptides useful for treating aids
TI
         Bolognesi, Dani Paul, Durham, NC, UNITED STATES
Matthews, Thomas James, Durham, NC, UNITED STATES
Wild, Carl T., Durham, NC, UNITED STATES
IN
         Barney, Shawn O?apos, Lin, Cary, NC, UNITED STATES
         Lambert, Dennis Michael, Cary, NC, UNITED STATES
         Petteway, Stephen Robert, Cary, NC, UNITED STATES
         Langlois, Alphonse J., Durham, NC, UNITED STATES
PA
         Duke University (U.S. corporation)
         Trimeris, Inc. (U.S. corporation) US 2004052820 A1 20040318
PΙ
         US 2004052820 A1 20040518
US 2002-267748 A1 20021008 (10)
Continuation of Ser. No. US 1995-484223, filed on 7 Jun 1995, PENDING
Division of Ser. No. US 1995-470896, filed on 6 Jun 1995, GRANTED, Pat.
No. US 6479055 Continuation-in-part of Ser. No. US 1994-360107, filed on
20 Dec 1994, GRANTED, Pat. No. US 6017536 Continuation-in-part of Ser.
No. US 1994-255208, filed on 7 Jun 1994, GRANTED, Pat. No. US 6440656
Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993,
ΑI
RLI
         GRANTED, Pat. No. US 5464933
DT
         Utility
FS
         APPLICATION
LN.CNT 40442
INCL
         INCLM: 424/208.100
         INCLS: 424/188.100; 530/350.000; 424/204.100; 530/300.000
NCL
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         NCLM:
                  424/188.100; 530/350.000; 424/204.100; 530/300.000
         NCLS:
IC
         [7]
         ICM: A61K039-21
         ICS: C07K014-16; A61K039-12; C07K002-00; C07K004-00; C07K005-00;
         C07K007-00; C07K014-00; C07K016-00; C07K017-00; A61K038-00; C07K001-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
      ANSWER 7 OF 76 USPATFULL on STN
AN
         2004:69539 USPATFULL
ΤI
         Immunization against amyloid plaques using display technology
         Solomon, Beka, Herzlia Pituach, ISRAEL
IN
         Frenkel, Dan, Rehovot, ISRAEL
         Ramot at Tel-Aviv University, Ltd., Tel Aviv, ISRAEL (non-U.S.
PA
         corporation)
PΙ
         us 2004052766
                                 Α1
                                        20040318
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         us 2003-618856
                                 Α1
                                        20030715 (10)
         Continuation of Ser. No. US 1999-473653, filed on 29 Dec 1999, ABANDONED
RLI
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PRAI
        US 1999-152417P
                              19990903 (60)
DT
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        APPLICATION
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NCL
        NCLM: 424/093.200
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IC
        ICM: A61K048-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 8 OF 76 USPATFULL on STN
        2004:44245
ΑN
                    USPATFULL
        Nucleic acids encoding DP-178 and other viral fusion inhibitor peptides
TI
        useful for treating aids
       Bolognesi, Dani Paul, Durham, NC, UNITED STATES
Matthews, Thomas James, Durham, NC, UNITED STATES
Wild, Carl T., Durham, NC, UNITED STATES
IN
        Duke University (U.S. corporation)
PA
PΙ
        us 2004033235
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                                  20040219
        us 2003-267682
ΑI
                             Α1
                                  20030106 (10)
        Continuation of Ser. No. US 1995-484223, filed on 7 Jun 1995, PENDING
RLI
        Division of Ser. No. US 1995-470896, filed on 6 Jun 1995, GRANTED, Pat.
        No. US 6479055 Continuation-in-part of Ser. No. US 1994-360107, filed on
        20 Dec 1994, GRANTED, Pat. No. US 6017536 Continuation-in-part of Ser. No. US 1994-255208, filed on 7 Jun 1994, GRANTED, Pat. No. US 6440656
        Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993,
        GRANTED, Pat. No. US 5464933
DT
        Utility
FS
        APPLICATION
LN.CNT 59510
        INCLM: 424/186.100
INCL
        INCLS: 424/188.100; 530/350.000; 424/208.100; 424/187.100
NCL
        NCLM:
               424/186.100
        NCLS:
               424/188.100; 530/350.000; 424/208.100; 424/187.100
IC
        [7]
        ICM: A61K039-21
        ICS: A61K039-12; C07K014-16; C07K014-10; C07K014-05; C07K014-11
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 9 OF 76 USPATFULL ON STN
L3
AN
        2004:181084 USPATFULL
ΤI
        Immunological detection of prions
IN
        Korth, Carsten, San Francisco, CA, United States
        Stierli, Beat, Daenikon, SWITZERLAND
       Stregt, Peter, Zurich, SWITZERLAND Oesch, Bruno, Stilli, SWITZERLAND
        Moser, Markus, Zurich, SWITZERLAND
PA
        Universitat Zurich, Zurich, SWITZERLAND (non-U.S. corporation)
PΙ
        us 6765088
                                  20040720
                             В1
        wo 9837210
                     19980827
                                  19990823 (9)
ΑI
        US 1999-380015
        WO 1998-EP917
                                  19980218
PRAI
        DE 1997-102837
                              19970221
        Utility
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LN.CNT
       1703
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        INCLM: 530/388.100
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        INCLS:
               530/388.850
               530/388.100
NCL
       NCLM:
               424/139.100; 424/141.100; 435/007.100; 435/326.000; 435/331.000;
        NCLS:
               530/388.850
IC
        [7]
        ICM: C07K016-00
EXF
        435/7.1; 435/326; 435/331; 436/503; 436/518; 436/547; 530/387.1;
        530/387.9; 530/388.1; 530/388.85; 800/4; 800/5; 800/6
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 10 OF 76 WPIDS COPYRIGHT 2004 THOMSON DERWENT ON STN
L3
     2004-501381 [48]
ΑN
                          WPIDS
DNN
     N2004-395892
                          DNC C2004-185869
     Amplifying immunodetection of pathological prion protein, useful for
ΤI
     diagnosis of spongiform encephalopathy, by adding a macrocyclic
        ***adjuvant***
                          ligand before reaction with antibody.
DC
     A26 A89 B04 D16 E19 S03
     COLEMAN, A W; DA SILVA, E; MARTIN, A; MOUSSA, A; SHAHGALDIAN, P; DUPIN, M;
IN
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LAZAR, A N: LECLERE, E: PERRON, H
     (FRSE-N) AGENCE FR SECURITE SANITAIRES ALIMENTS; (CNRS) CNRS CENT NAT RECH
PA
     SCI; (UYLY-N) UNIV LYON 1 BERNARD CLAUDE; (INMR) BIOMERIEUX SA
CYC
     107
                      A1 20040625 (200448)*
A1 20040715 (200448)
     FR 2849205
PΙ
                                                         G01N033-68
     wo 2004059322
                                            FR
                                                         G01N033-68
        RW: AT BE BG BW CH CY CZ DE DK EA EE ES FI FR GB GH GM GR HU IE IT KE
             LS LU MC MW MZ NL OA PT RO SD SE SI SK SL SZ TR TZ UG ZM ZW
         W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK
             DM DZ EC EE EG ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP
             KR KZ LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NI NO NZ OM PG
             PH PL PT RO RU SC SD SE SG SK SL SY TJ TM TN TR TT TZ UA UG US UZ
             VC VN YU ZA ZM ZW
     FR 2849205 A1 FR 2002-16383 20021220; WO 2004059322 A1 WO 2003-FR3857
ADT
     20031219
PRAI FR 2002-16383
                           20021220
          G01N033-68
IC
     ICM
          C12Q001-37; G01N033-566
     ICS
      ANSWER 11 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT ON STN
L3
ΑN
      2004-27090 DRUGU
                           T S
TI
      Thiotepa-based high-dose chemotherapy with autologous stem-cell rescue in
      patients with recurrent or progressive CNS germ cell tumors.
ΑU
      Modak S; Gardner S; Dunkel I J; Balmaceda C; Rosenblum M K; Miller D C;
      Halpern S; Finlay J L
      Mem.Sloan-Kettering-Cancer-Cent.; Univ.New-York-Columbia; Univ.New-York
      New York, N.Y., USĂ
J.Clin.Oncol. (22, No. 10, 1934-43, 2004) 3 Fig. 5 Tab. 40 Ref.
LO
S0
      CODEN: JCONDN
                           ISSN:
                                  0732-183X
ΑV
      Department of Pediatrics, Memorial Sloan-Kettering Cancer Center, 1275
      York Ave, New York, NY 10021, U.S.A. (e-mail: modaks@mskcc.org).
LA
      English
DT
      Journal
      AB; LA; CT
FΑ
FS
      Literature
L3
     ANSWER 12 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation. on
     STN
                                                           DUPLICATE 2
     2004:365298 BIOSIS
AN
DN
     PREV200400368764
     Multiple antigenic peptides facilitate generation of anti-prion
TI
     antibodies.
     Bainbridge, J.; Jones, N.; Walker, B. [Reprint Author]
ΑU
CS
     Dept Immunobiol, Natl Inst Biol Stand and Controls, Blanche Lane S Mimms,
     Potters Bar, Herts, EN6 3QG, England
     kbwalker@nibsc.ac.uk
50
     Clinical and Experimental Immunology, (August 2004) Vol. 137, No. 2, pp.
     298-304. print.
     ISSN: 0009-9104 (ISSN print).
DT
     Article
LA
     Enalish
ED
     Entered STN: 8 Sep 2004
     Last Updated on STN: 8 Sep 2004
L3
     ANSWER 13 OF 76 CAPLUS COPYRIGHT 2004 ACS ON STN DUPLICATE 3
ΑN
     2003:930741 CAPLUS
DN
     140:4064
     vaccine comprising conjugates of prion protein and recombinant virus-like
TI
     particle carrier for treating prion diseases
     Bachmann, Martin; Maurer, Paťrik; Pellicioli, Erica; Renner, Wolfgang A.
IN
PA
     Cytos Biotechnology Ag, Switz.
     U.S. Pat. Appl. Publ., 127 pp., Cont.-in-part of Appl. No. PCT/IB02/00166.
SO
     CODEN: USXXCO
DT
     Patent
     English
LA
FAN.CNT 9
     PATENT NO.
                          KIND
                                 DATE
                                              APPLICATION NO.
                                                                       DATE
PΙ
     US 2003219459
                           Α1
                                  20031127
                                              US 2003-346190
                                                                       20030117
     US 2003175290
                                  20030918
                                              US 2002-50902
                           Α1
                                                                       20020118
     wo 2002056905
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                                  20020725
                                              WO 2002-IB166
                                                                       20020121
     wo 2002056905
                           Α3
                                 20031009
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             CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, ES, FI, GB, GD, GE, GH,
             GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, OM, PH,
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PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TN, TR, TT, TZ,
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                   TJ, TM
             RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZM, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG 2002-50902 A2 20020118
PRAI US 2002-50902
       WO 2002-IB166
                                      Α2
                                               20020121
       US 2002-393725P
                                               20020708
       US 2002-396590P
                                               20020718
       US 2001-262379P
                                      Ρ
                                               20010119
       US 2001-288549P
                                      Р
                                               20010504
       US 2001-326998P
                                      Ρ
                                               20011005
       US 2001-331045P
                                               20011107
       ANSWER 14 OF 76 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 4
L3
         10422134 IFIPAT; IFIUDB; IFICDB
ΑN
         SYNTHETIC IMMUNOGENIC BUT NON-DEPOSIT-FORMING POLYPEPTIDES AND PEPTIDES
ΤI
         HOMOLOGOUS TO AMYLOID BETA, PRION PROTEIN, AMYLIN, ALPHA-SYNUCLEIN, OR
         POLYGLUTAMINE REPEATS FOR INDUCTION OF AN IMMUNE RESPONSE THERETO
IN
         Frangione Blas; Sigurdsson Einar M; Wisniewski Thomas
PA
         New York University (59449)
PΙ
         us 2003166558
                               Α1
                                      20030904
         US 2002-301488
                                      20021121
AΤ
         US 2001-331801P
PRAI
                                      20011121 (Provisional)
FI
         US 2003166558
                                      20030904
         Utility; Patent Application - First Publication
DT
FS
         CHEMICAL
         APPLICATION
CLMN
         115
GΙ
          15 Figure(s).
       FIG. 1 shows the results of a thioflavin T fluorometric assay. Fibril
         formation of A beta 1-42, A beta 1-30-NH2, and K6A beta 1-30-NH2 (SEQ ID
         NO:6) was measured in vitro following incubation at 37 degrees C. K6A
         beta 1-30-NH2 was the only peptide that did not form fibrils at any of
         the time points.
       FIGS. 2A and 2B show that A beta 40 and A beta 42 are toxic to human
        neuroblastoma cells (SK-N-SH) in culture as determined by the MTT assay, whereas K6A beta 30-NH2 has no effect at 2 days (FIG. 2A) and is slightly trophic at 6 days (FIG. 2B). *p lessthan 0.05; **p less-than 0.01; ***p
       less-than 0.001 compared to VEH group (one-way ANOVA). FIGS. 3A-3D show coronal sections (X50; original magnification) stained
        with 6E10 against A beta, through the hippocampus and cortex in a Tg control-(FIG. 3A) and K6A beta 1-30-treated (FIG. 3B) Tg mouse. FIGS. 3C
         and 3D are adjacent sections (X100) double stained for interleukin-1 that
         recognizes microglia, and A beta . Note the reduction of amyloid burden in the immunized mouse (FIG. 3B), and the lack of ramified microglia
        (FIG. 3D) surrounding A beta plaque in the same mouse, compared to a control mouse (FIG. 3A, 3C). The bars in FIGS. 3A and 3C are 100 mu m.
       Abbreviations: hip=hippocampus; cx=cortex; cc=corpus callosum. FIGS. 4A-4C show the reduction in cortical (FIG. 4A) and hippocampal (FIG.
         4B) amyloid burden (6E10) following 7 months treatment with K6A beta
         1-30-NH2. There is an 89% reduction in cortical amyloid burden
        (*p=0.0002; t-test; n=4 per group) and an 81% reduction in hippocampal amyloid burden (*p=0.0001). Soluble A beta 1-42 levels (FIG. 4C) are reduced by 57% within the brains of the vaccinated mice (*p=0.0019).
       FIG. 5 shows the results of a thioflavin T fluorometric assay. Fibril
         formation of A beta 1-42, A beta 1-40, A beta 1-30-NH2, A beta 1-30K6, A
        beta 1-30-NH2 (EE18,19) and A beta 1-30-NH2 (DDL18,19) was measured in vitro following incubation at 37 degrees C. for 15 days. Within this
         period, no fibril formation of the A beta derivatives containing a
         polylysine segment or an amino acid substitution within the hydrophobic
         region was detected.
       FIGS. 6A and 6B show the results of MTT cell toxicity assay. Neurotoxicity
        of A beta 1-42, A beta 1-40, A beta 1-30-NH2, K6A beta 1-30-NH2, A beta 1-30K6, A beta 1-30-NH2(EE,18,19) and A beta 1-30-NH2(DD,18,19) was determined following treatment of human neuroblastoma cells (SK-N-SH) for 2 (FIG. 6A) and 6 (FIG. 6B) days. *p less-than 0.05; **p less-than 0.01; ***p less-than 0.001 compared to VEH group (one-way ANOVA). In this*** assay, A beta 1-40 and A beta 1-42 were toxic to human neuroblastoma***
  re re re
                cells (SK-N-SH) in culture. Of the A beta derivatives, even at the*** highest concentration (100 mu M), only A beta 1-30K6 displayed a slight***
   ** ** **
                toxicity and only on day 2 of the test. Several of the peptides were neurotrophic following 6 days incubation. *p less-than 0.05; **p***
                less-than 0.01; ***p lessthan 0.001 (One-way Anova; Neuman Keuls' posthoc***
                test). ***
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FIG. 7 shows the antibody titer determined by ELISA in mice 14 weeks after***
  de de de
  te te te
             vaccination with mouse recPrP. ***
  de de de
            FIGS. 8A and 8B show that a higher anti-PrPC (ME7 FAS PrP) antibody titer***
             in vaccinated mice, as presented in FIG. 7, correlates with a longer***
  de de de
                                            ***PrPSc***
             incubation time in both
                                                              inoculated mouse groups at lower
       dilution (FIG. 8A; r2=0.4389, p=0.0052) and at higher dilution (FIG. 8B;
       r2=0.6786, p lessthan 0.0001)
      FIG. 9 is a graph showing the effect of recPrP vaccination on disease onset, with day 0 being the first day an animal scored positive for
       disease. Group 1 mice were controls inoculated with
                                                                          ***PrPSc***
       10 fold dilution, while group 2 was inoculated at the same dilution but
       also received recPrP vaccination. Group 3 mice were controls inoculated
       with ***PrPSc*** at a 1000 fold dilution, while Group 4 received the same dilution of ***PrPSc*** along with recPrP vaccination. The two
       control groups received ***adjuvant***
                                                           and vehicle injections. Two
       way ANOVA shows a significant effect for vaccination (p=0.0005) and
          ***PrPSc*** dilution (p less-than 0.000001). The Newman-Keuls post-hoc
       test showed vaccination to have a stronger effect in the 10 fold dilution
       group (Group 1 versus 2, p=0.001 two-tailed; Group 3 versus 4, p=0.036
       one-tailed).
      FIG. 10 shows an alignment of amino acid sequences of prion protein (PrP)
       from human (SEQ ID NO:21), gorilla (SEQ ID NO:22), chimpanzee (SEQ ID NO:23), mouse (SEQ ID NO:24), rat (SEQ ID NO:25), Syrian hamster (SEQ ID NO:26), might (SEQ ID NO:27), strate (SEQ ID NO:27)
       NO:26), mink (SEQ ID NO:27), sheep (SEQ ID NO:28), goat (SEQ ID NO:29),
       cow (SEQ ID NO:30), and greater kudu (SEQ ID NO:31). Amino acid residues
       that are identical and conserved among the prion proteins of the species
       presented in this figure are boxed.
      FIGS. 11A-C show ELISA evaluation of sera from individual animals vaccinated with K6A beta 1-30-NH2 and alum ***adjuvant***, t
       for antibody titer against antigen (FIG. 11A), A beta 142 (FIG. 11B) and
       A beta 1-40 (FIG. 11C).
      FIGS. 12A-C show ELISA evaluation of sera from individual animals immunized with A beta 1-42 and alum ***adjuvant***, testing for
       antibody titer against antigen (FIG. 12A), K6AP1-30-NH2 (FIG. 12B) and A
       beta 1-40 (FIG. 12C)
      FIGS. 13A and 13B depict a linear maze used to evaluate cognitive
       capabilities of animals vaccinated with A beta 1-30NH2 and K6A beta
       1-30-NH2 together with alum ***adjuvants*** , as well as controls.
      FIG. 13A shows the maze design during the adaptation phase, and FIG. 13B during testing. Dotted lines indicate blocked alleys.
      FIGS. 14A-C depict results obtained from behavioral studies of animals of
       about 3-4 months of age, after vaccination with A beta 1-30-NH2 and K6A
       beta 1-30-NH2 together with alum ***adjuvants*** , as well as
       controls. The studies included testing of locomotor activity (FIG. 14A),
       spontaneous avoidance (FIG. 14B), and passive avoidance (FIG. 14C). See
       Example 6.
      FIGS. 15A-N depict results obtained from behavioral studies of animals of
       about 11 months of age, after vaccination with A beta 1-30-NH2 and K6A beta 1-30-NH2 together with alum ***adjuvants***, as well as
       beta 1-30-NH2 together with alum ***adjuvants*** , as well as controls. The studies included testing of locomotor activity (FIG. 15A), and cognitive testing using traverse beam (FIGS. 15B and 15C), rotarod (FIG. 15D), radial arm maze (FIGS. 15E and 15F), straight alley channel (FIGS. 15G), visible platform (FIGS. 15H and 15T), Morris water maze (FIGS. 15L and 15V)
       (FIGS. 15J and 15K), probe trial (FIGS. 15L and 15M), and linear maze
       (FIG. 15N). See Example 6.
L3
      ANSWER 15 OF 76 IFIPAT COPYRIGHT 2004 IFI on STN DUPLICATE 5
AN
       10304838 IFIPAT; IFIUDB; IFICDB
       AGENT; THERAPY, PREVENTION PRION INFECTIONS; USING MONOCLONAL ANTIBODIES ENCAPSULATED AS HYBRIDOMAS
TI
       Enari Masato (JP); Weissmann Charles (GB)
TN
PA
       Unassigned Or Assigned To Individual (68000)
PΙ
       US 2003049249
                         A1 20030313
ΑI
       US 2001-985164
                                20011101
PRAI
       GB 2001-221621
                                20010913
FΙ
       US 2003049249
                                20030313
DT
       Utility; Patent Application - First Publication
FS
       CHEMICAL
       APPLICATION
os
       CA 138:231762
CLMN
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        4 Figure(s).
      FIG. 1 represents the susceptibility to scrapie infection and PrP level of
       various sublines of N2a cells. N2a populations as propagated routinely in
       the lab and single clones transformed with a PrP expression or a control
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vector are seeded into 24well plates (2 x 104 cells/well) and grown to

confluence. (a) Cultures are exposed for 3 days to purified mouse

***PrPSc*** (RML strain, 20 ng/ml), cultured for 29 days (8 passages)
and assayed for ***PrPSc*** formation by the cell blot assay. (b) Prionsusceptible N2a/Bos2 and resistant N2a/2M11 cells are exposed for 3 days to the dilutions indicated of infected 10% brain homogenate, cultured for 14 days (3 passages) and assayed for ***PrPSC*** cultured for 14 days (3 passages) and assayed for ***PrPSc*** formation. Cells exposed to a 10-4 dilution are still slightly positive (c) Western blot analysis of N2a sublines is performed using monoclonal anti-PrP antibody 6H4. Cells transfected with the expression plasmid for mouse PrPc, MHM2 PrP or MH2M PrP are indicated by mo, M2 or 2M, respectively. BOS designates cells cotransfected with pSVneo and pEF-BOS-EX. N2a, the original uncloned cells, as well as the highly susceptible N2a/Bos2 cell line show similar, low expression of PrPC as compared to the non-susceptible mos or 2M11 lines. FIG. 2 represents anti PrP antibody 6H4 and PIPLC preventing infection of N2a/Bos2 cell with scrapie prions and abrogate ***PrPSc*** N2a/Bos2 cell with scrapie prions and abrogate accumulation in chronically infected cells. (a) N2a/Bos2 cells are incubated for 2 h with antibody 6H4 or PIPLC at the concentrations indicated and exposed to 0.1% scrapie-infected brain homogenate (fmal concentration) for 3 days. After culturing for 14 days (3 passages) in the absence of PIPLC or in the continued presence of 6H4, ***PrPSC**** expression is determined. (b,c) Chronically scrapie-infected N2a/Bos2 cells are cultured for (b) 3 days at the levels of antibody 6H4 or of PIPLC indicated or (c) 14 days (3 passages) at the concentrations of antibody 6H4 indicated, and ***PrPSc*** accumulation is monitored. (d) Chronically scrapie-infected N2a/ Bos2 cells are exposed to 6H4 at the concentrations indicated for 2 weeks and further cultured in the absence of the antibody for 6 weeks. Cultures are split 1:5 every 34 days. There is no reappearance of ***PrPSc*** . "Cell staining" refers to staining of the membranes with ethidium bromide to monitor efficiency of transfer of the cell layer. IN, chronically scrapie-infected N2a/Bos2 FIG. 3 represents chronically infected N2a/Bos2 cells "cured" of ***PrPSc*** by antibody 6H4 treatment continuing to produce PrP and their susceptibility to reinfection. (a) Chronically infected N2a/Bos2 cells, treated with antibody 6H4 for 2 weeks at the concentrations indicated and propagated in its absence for 66 days were not exposed (top 2 rows) or exposed to 0.1% RMLinfected mouse brain homogenate for 3 days (hottom 2 rows). After culturing for 14 days, ***PrPSc*** is (bottom 2 rows). After culturing for 14 days, ***PrPSc*** is monitored by the cell blot assay. (b) Relative susceptibility to prions of N2a/Bos2 cells (Bos2) and N2a/Bos2 cells "cured" by exposure to antibody 6H4 at 20 mu g/ml is determined by exposing cultures to various dilutions of RML-infected mouse brain homogenate for 3 days and ***PrPSc*** as above. (c) Levels of PrPC and in various sublines are determined by Western blotting. Chronically infected N2a/Bos2 cells, treated for 2 weeks with antibody 6H4 at the concentrations indicated are passaged for 84 days after antibody withdrawal. Cells are lysed and samples corresponding to 2.25 x 105 cells are incubated in the presence (+PK) or absence (-PK) of proteinase K (5 mu g/ml) for 90 minutes at 37 degrees C. Western blotting is performed as described in Methods. UN, uninfected N2a/Bos2 cells and I-BOS2, chronically prion-infected BOS2 cells. Molecular weight markers are indicated at the left of each panel ***PrPSc*** FIG. 4 represents a model to explain abolition of bv anti PrP antibody (or PIPLC). PrPC is attached to the membrane by a glycosylphosphatidyl inositol anchor and cycles between the cell surface and an endocytic compartment (43). In scrapie infected cells, PrPC is recruited into ***PrPSc*** "seeds" (44), which may be located at the cell surface and/or in the endocytic/lysosomal compartment. ***PrPSc*** is degraded with a halflife of about 15 h (37); if PrPC is prevented from converting to ***PrPSc*** by either a blocking antibody or by being stripped from the cell surface by PIPLC, ***PrPSc*** will diminish and ultimately disappear. ANSWER 16 OF 76 CAPLUS COPYRIGHT 2004 ACS ON STN

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AN 2004:2727 CAPLUS
DN 140:75940
TI Unmethylated CpG oligonucleotide-packaged virus-like particles for enhancing immune response of vaccines
IN Bachman, Martin F.; Renner, Wolfgang A.
PA Cytos Biotechnology Ag, Switz.
SO PCT Int. Appl., 252 pp.
CODEN: PIXXD2
DT Patent
LA English
FAN.CNT 1
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PRAI US 2002-389898P
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                 THERE ARE 5 CITED REFERENCES AVAILABLE FOR THIS RECORD
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AN
DN
      139:128014
      Methods for treating diseases or conditions with peptide constructs
TI
      Zimmerman, Daniel H.; Charoenvit, Yupin; Rosenthal, Kenneth S.; Whelan,
IN
      Mike
PA
      Cel-Sci Corporation, USA
      PCT Int. Appl., 73 pp.
S0
      CODEN: PIXXD2
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      English
LA
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PRAI US 2002-349982P P 2002
                                       20020123
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                               Ρ
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                               Ρ
                                       20020123
      ANSWER 18 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
L3
AN
      2003:570846
                    CAPLUS
DN
      139:132445
TI
      Pharmaccine comprising conjugates of prion protein and virus-like particle
      carrier for treating prion diseases
ΙN
      Bachmann, Martin; Maurer, Patrick; Pellicioli, Erica; Renner, Wolfgang A.
      Cytos Biotechnology A.-G., Switz.
PA
S0
      PCT Int. Appl., 247 pp.
      CODEN: PIXXD2
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LA
FAN.CNT 9
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PRAI US 2002-50902
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     US 2002-396590P
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                                    20010119
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                                    20011005
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                             Ρ
                                    20011107
L3
     ANSWER 19 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
      2003:434266 CAPLUS
ΑN
      139:21013
DN
      Synthetic immunogenic/non-deposit-forming polypeptides and peptides
ΤI
     hómologous to amyloid .beta., prion protein, amylin, .alpha.-synuclein, or polyglutamine repeats for induction of an immune response
     Frangione, Blas; Wisniewski, Thomas; Sigurdsson, Einar M.
ΙN
     New York University, USA
PA
S0
     PCT Int. Appl., 265 pp.
     CODEN: PIXXD2
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PRAI US 2001-331801P
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L3
     ANSWER 20 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
     2003:334937 CAPLUS
ΑN
DN
     138:349696
     Fusion protein comprising cytokines, chemokines, and interferons for use as vaccine ***adjuvant*** in immunotherapy for cancer and viral
TI
     infection
     Galipeau, Jacques; Stagg, John
IN
     Centre for Translational Research In Cancer, Can.
PA
     PCT Int. Appl., 53 pp.
SO
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PRAI US 2001-330476P
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     ANSWER 21 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
     2003:242184 CAPLUS
AN
DN
     138:285995
     Packaging of immunostimulatory substances and antigens into virus-like
TI
     particles for use as vaccines against cancer, autoimmune disease, allergy
     and viral infection
     Maurer, Patrick; Tissot, Alain; Schwarz, Katrin; Meijerink, Edwin; Lipowsky, Gerad; Pumpens, Paul; Cielens, Indulis; Renhofa, Regina;
ΙN
     Bachmann, Martin F.; Storni, Tazio
     Cytos Biotechnology A.-G., Switz.
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     PCT Int. Appl., 322 pp.
     CODEN: PIXXD2
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     ANSWER 22 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
     2003:242183 CAPLUS
ΑN
DN
     138:270293
TI
     Vaccine compositions comprising anti-CD4 antibody or immunostimulatory
     nucleic acid and antigen-coupled virus-like particles for enhancement of
     immune responses
IN
     Bachmann, Martin F.; Storni, Tazio; Lechner, Franziska
     Cytos Biotechnology A.-G., Switz.
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SO
     PCT Int. Appl., 243 pp.
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     ANSWER 23 OF 76 IFIPAT COPYRIGHT 2004 IFI on STN
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      Fishleigh Robert Vincent (GB); Mee Roger Paul (GB); Robson Barry (GB)
      Proteus Molecular Design Ltd GB (43521)
PA
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      US 2002-116061
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      Utility; Patent Application - First Publication
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L3
     ANSWER 24 OF 76 USPATFULL ON STN
       2003:334947 USPATFULL
AN
       Nucleotide sequences that code for torsin genes, torsin proteins, and
TI
       methods of using the same to treat protein-aggregation
       Caldwell, Guy A., Tuscaloosa, AL, UNITED STATES
IN
       Caldwell, Kim A., Tuscaloosa, AL, UNITED STATES
       THE UNIVERSITY OF ALABAMA, Tuscaloosa, AL (U.S. corporation)
PA
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       us 2003235823
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       APPLICATION
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               530/388.100; 536/023.500; 514/012.000
NCL
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               530/388.100; 536/023.500; 514/012.000
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       ICM: C12Q001-68
       ICS: G01N033-53; C12P021-02; C12N005-06; C07K014-47; C07K016-20;
       A61K038-17
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 25 OF 76 USPATFULL ON STN
ΑN
       2003:330543
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TT
       Immunological methods and compositions for the treatment of Alzheimer's
       disease
       St. George-Hyslop, Peter H., Toronto, CANADA
McLaurin, JoAnne, Toronto, CANADA
Hospital for Sick Children and University of Toronto (non-U.S.
IN
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       corporation)
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NCL
               514/012.000
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               530/324.000; 435/069.100; 435/320.100; 435/325.000; 536/023.100
       NCLS:
       [7]
       ICM: A61K038-17
       ICS: C07K014-47; C12P021-02; C12N005-06
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 26 OF 76 USPATFULL ON STN
```

```
2003:324329 USPATFULL
        Antibodies specific for native
                                           ***PrPsc***
TI
IN
        Prusiner, Stanley B., San Francisco, CA, UNITED STATES
        Williamson, R. Anthony, San Diego, CA, UNITED STATES
Burton, Dennis R., LaJolla, CA, UNITED STATES
The Regents of the University of California and The Scripps Research
PA
        Institute (U.S. corporation)
        us 2003228303
PΙ
                            A1
                                  20031211
        US 2003-435602
ΑI
                            Α1
                                  20030509 (10)
        Continuation of Ser. No. US 2001-943906, filed on 30 Aug 2001, GRANTED,
RLI
        Pat. No. US 6562341 Continuation of Ser. No. US 2000-550374, filed on 13
        Apr 2000, GRANTED, Pat. No. US 6372214 Continuation of Ser. No. US
        1998-36579, filed on 6 Mar 1998, GRANTED, Pat. No. US 6290954 Division
        of Ser. No. US 1996-713939, filed on 13 Sep 1996, GRANTED, Pat. No. US
        5846533 Continuation-in-part of Ser. No. US 1995-528104, filed on 14 Sep
        1995, ABANDONED
DΤ
        Utility
        APPLICATION
FS
LN.CNT 2983
INCL
        INCLM: 424/130.100
NCL
        NCLM: 424/130.100
IC
        [7]
        ICM: A61K039-395
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 27 OF 76 USPATFULL ON STN
        2003:264865 USPATFULL
ΑN
        Therapy for human cancers using cisplatin and other drugs or genes
TI
        encapsulated into liposomes
ΙN
        Boulikas, Teni, Palo Alto, CA, UNITED STATES
        US 2003185879
PΙ
                           Α1
                                 20031002
ΑI
        US 2003-350470
                            Α1
                                 20030123 (10)
       Division of Ser. No. US 1999-434345, filed on 5 Nov 1999, GRANTED, Pat.
RLI
       No. US 6511676
DT
        Utility
        APPLICATION
FS
LN.CNT
       1652
INCL
        INCLM: 424/450.000
        INCLS: 424/649.000
               424/450.000
NCL
       NCLM:
        NCLS:
               424/649.000
IC
        [7]
        ICM: A61K009-127
        ICS: A61K033-24
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 28 OF 76 USPATFULL ON STN
ΑN
       2003:264794 USPATFULL
       Prostate cell lines
TT
IN
       Thraves, Peter, London, UNITED KINGDOM
       Sutton, Andrew, London, UNITED KINGDOM
PΙ
       US 2003185808
                            Α1
                                 20031002
ΑI
       US 2002-240523
                                 20021023 (10)
                            Α1
       WO 2001-GB1437
                                 20010330
PRAI
       GB 2000-8032
                             20000401
       GB 2000-24237
                             20001003
       Utility
DT
       APPLICATION
FS
LN.CNT 727
INCL
       INCLM: 424/093.210
       INCLS: 514/044.000; 435/366.000; 424/085.200
NCL
               424/093.210
       NCLS:
               514/044.000; 435/366.000; 424/085.200
IC
       [7]
       ICM: A61K048-00
       ICS: C12N005-08
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 29 OF 76 USPATFULL ON STN
L3
       2003:206867 USPATFULL
ΑN
       Antibodies specific for ungulate PrP
TI
IN
       Prusiner, Stanley B., San Francisco, CA, UNITED STATES
       Safar, Jiri G., Walnut Creek, CA, UNITED STATES
       Williamson, R. Anthony, San Diego, CA, UNITED STATES
       Burton, Dennis R., La Jolla, CA, UNITED STATES
PΙ
       US 2003143224
                            Α1
                                 20030731
```

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ΑI
        us 2003-355780
                                 20030130 (10)
                            Α1
        Continuation of Ser. No. US 2000-627218, filed on 27 Jul 2000, GRANTED,
RLI
        Pat. No. US 6537548
DT
        Utility
FS
        APPLICATION
LN.CNT 2123
        INCLM: 424/130.100
INCL
        INCLS: 435/345.000; 435/006.000
        NCLM: 424/130.100
NCL
        NCLS:
               435/345.000; 435/006.000
        [7]
IC
        ICM: C12Q001-68
        ICS: A61K039-395; C12N005-06; C12N005-16
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
      ANSWER 30 OF 76 USPATFULL ON STN
        2003:181684
ΑN
                     USPATFULL
        Surface simulation synthetic peptides useful in the treatment of
ΤI
        hyper-variable viral pathogens
IN
        Crevecoeur, Harry F., Valley Stream, NY, UNITED STATES
PΙ
        US 2003125518
                                 20030703
                            Α1
        us 2001-12806
ΑI
                            Α1
                                 20011201 (10)
DT
        Utility
FS
        APPLICATION
LN.CNT 1650
INCL
        INCLM: 530/327.000
               530/328.000; 435/005.000; 702/019.000
               530/327.000
NCL
        NCLM:
        NCLS:
               530/328.000; 435/005.000; 702/019.000
IC
        [7]
        ICM: C07K007-08
        ICS: C12Q001-70; G01N033-48; G06F019-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 31 OF 76 USPATFULL ON STN
        2003:126727 USPATFULL
ΑN
TI
       Novel methods for down-regulation of amyloid
TN
       Jensen, Martin Roland, Horsholm, DENMARK
       Birk, Peter, Horsholm, DENMARK
       Nielsen, Klaus Gregorius, Horsholm, DENMARK
ΡI
       US 2003086938
                                 20030508
                            Α1
ΑI
       US 2002-204362
                            Α1
                                 20020816 (10)
       WO 2001-DK113
                                 20010219
PRAI
       DK 2000-265
                             20000221
       Utility
DT
       APPLICATION
FS
LN.CNT
       3114
INCL
       INCLM: 424/185.100
NCL
       NCLM:
             424/185.100
IC
        [7]
       ICM: A61K039-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 32 OF 76 USPATFULL ON STN
       2003:112522 USPATFULL
ΑN
TI
       Agents and compositions and methods utilizing same useful in diagnosing
       and/or treating or preventing plaque forming
       Solomon, Beka, Herzlia Pituach, ISRAEL
Hanan, Eilat, Tel Aviv, ISRAEL
Frenkel, Dan, Rehovot, ISRAEL
ΙN
       Ramot University Authority for Applied Research & Industrial
PA
       Development, Tel Aviv, ISRAEL (non-U.S. corporation)
PΙ
       US 2003077252
                           Α1
                                 20030424
ΑI
       US 2002-162889
                           Α1
                                 20020606 (10)
       Continuation of Ser. No. US 2000-629971, filed on 31 Jul 2000, ABANDONED
RLI
       Continuation-in-part of Ser. No. US 1999-473653, filed on 29 Dec 1999,
       PENDING
PRAI
       US 1999-152417P
                            19990903 (60)
DT
       Utility
       APPLICATION
FS
LN.CNT 2994
INCL
       INCLM: 424/093.200
       INCLS: 514/044.000; 435/456.000; 435/235.100
NCL
       NCLM:
              424/093.200
       NCLS:
              514/044.000; 435/456.000; 435/235.100
IC
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ICM: A61K048-00
        ICS: C12N007-01; C12N015-86
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 33 OF 76 USPATFULL ON STN
L3
        2003:44722 USPATFULL
AN
ΤI
        Early pre-symptomatic prion diagnostic blood test for encephalopathies
       Resink, Annelies, Paris, FRANCE
Fuentes, Nathalie, Kremlin Bicetre, FRANCE
IN
        Schweighoffer, Fabien, Vincennes, FRANCE
PI
        us 2003032032
                            Α1
                                  20030213
        us 2002-100178
                                  20020319 (10)
ΑI
                            Α1
        US 2001-278670P
                             20010321 (60)
PRAI
                             20010410 (60)
        US 2001-282463P
DT
        Utility
FS
        APPLICATION
LN.CNT
       962
        INCLM: 435/006.000
INCL
       NCLM: 435/006.000
NCL
IC
        [7]
        ICM: C12Q001-68
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 34 OF 76 USPATFULL on STN
        2003:273427 USPATFULL
AN
TT
        Inhibitors of amyloid formation
       Caughey, Winslow S., Hamilton, MT, United States Caughey, Byron, Hamilton, MT, United States
IN
       The United States of America as represented by the Department of Health
PA
       and Human Services, Washington, DC, United States (U.S. government)
                                  20031014
       US 6632808
                            В1
       wo 2000009111 20000224
       US 2001-762725
AΤ
                                  20010307 (9)
       wo 1999-us18297
                                  19990811
PRAI
       US 1998-96148P
                             19980811 (60)
       Utility
DT
FS
        GRANTED
LN.CNT
       1503
INCL
       INCLM: 514/185.000
       INCLS: 514/410.000; 540/122.000; 540/145.000
               514/185.000
NCL
       NCLM:
       NCLS:
               514/410.000; 540/122.000; 540/145.000
        [7]
IC
       ICM: A61K031-409
       ICS: C07D487-22
EXF
        514/185; 514/410; 540/122; 540/145
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 35 OF 76 USPATFULL ON STN
L3
ΑN
       2003:81453 USPATFULL
TI
       Antibodies specific for ungulate PrP
IN
       Prusiner, Stanley B., San Francisco, CA, United States
       Safar, Jiri, Concord, CA, United States
       Williamson, R. Anthony, San Diego, CA, United States
       Burton, Dennis R., La Jolla, CA, United States
The Regents of the University of California, Oakland, CA, United States
PA
        (U.S. corporation)
       The Scripps Research Institute, La Jolla, CA, United States (U.S.
       corporation)
PΙ
       US 6537548
                                  20030325
                            В1
       US 2000-627218
ΑI
                                  20000727 (9)
DT
       Utility
FS
       GRANTED
LN.CNT 2073
INCL
       INCLM: 424/130.100
       INCLS: 424/009.100; 424/185.100; 435/007.100; 435/070.100; 435/071.100;
               530/387.100; 530/398.100
NCL
               424/130.100
       NCLS:
               424/009.100; 424/185.100; 435/007.100; 435/070.100; 435/071.100;
               530/387.100; 530/389.100
       [7]
TC
       ICM: A61K039-395
       424/9.1; 424/130.1; 424/185.1; 435/7.1; 435/70.1; 435/71.1; 436/503;
EXF
       436/547; 530/387.1; 530/388.27; 530/398.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
```

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L3
      ANSWER 36 OF 76 USPATFULL on STN
        2003:40533 USPATFULL
AN
ΤI
        Methods for the inhibition of epstein-barr virus transmission employing
        anti-viral peptides capable of abrogating viral fusion and transmission
        Barney, Shawn O'Lin, Cary, NC, United States
Lambert, Dennis Michael, Cary, NC, United States
Petteway, Stephen Robert, Cary, NC, United States
Trimeris, Inc., Durham, NC, United States (U.S. corporation)
ΙN
PA
PI
        US 6518013
                                  20030211
                            В1
ΑI
        US 1995-485546
                                  19950607 (8)
        Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994,
RLI
        now patented, Pat. No. US 6017536 Continuation-in-part of Ser. No. US
        1994-255208, filed on 7 Jun 1994 Continuation-in-part of Ser. No. US
        1993-73028, filed on 7 Jun 1993, now patented, Pat. No. US 5464933
DT
        Utility
FS
        GRANTED
LN.CNT 24700
INCL
        INCLM: 435/005.000
        INCLS: 424/230.100; 530/300.000; 530/324.000; 530/325.000; 530/326.000
NCL
               435/005.000
               424/230.100; 530/300.000; 530/324.000; 530/325.000; 530/326.000
        NCLS:
IC
        [7]
        ICM: C12Q001-70
EXF
        435/5; 530/300; 530/324-329; 530/350; 424/230.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
1.3
     ANSWER 37 OF 76 USPATFULL ON STN
AN
        2003:26157
                    USPATFULL
        Therapy for human cancers using cisplatin and other drugs or genes
TI
        encapsulated into liposomes
ΙN
        Boulikas, Teni, 249 Matadero Ave., Palo Alto, CA, United States 94306
PΙ
        US 6511676
                            в1
                                  20030128
ΑI
        US 1999-434345
                                  19991105 (9)
        Utility
DT
FS
        GRANTED
LN.CNT
       1642
INCL
        INCLM: 424/450.000
        INCLS: 264/004.100; 264/004.300
NCL
        NCLM:
               424/450.000
        NCLS:
               264/004.100; 264/004.300
IC
        [7]
        ICM: A61K009-127
EXF
        424/450; 264/4.1; 264/4.3
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
      ANSWER 38 OF 76 DRUGU COPYRIGHT 2004 THOMSON DERWENT ON STN
      2004-07769 DRUGU
AN
TI
      Immunotherapy as a therapeutic treatment for neurodegenerative disorders.
ΑU
      White A R; Hawke S H
CS
      Howard-Florey-Inst.; Univ.London
      Melbourne, Austr.; London, U.K.
LO
SO
      J.Neurochem. (87, No. 4, 801-08, 2003) 1 Fig. 56 Ref.
      CODEN: JONRA9
                            ISSN: 0022-3042
      Neurochemistry Group, Howard Florey Institute of Experimental Physiology
ΑV
      and Medicine, Victoria 3010, Australia. (e-mail:
      a.white@hfi.unimelb.edu.au).
LA
      English
      Journal
DT
FΑ
      AB; LA; CT
FS
      Literature
L3
     ANSWER 39 OF 76 BIOSIS COPYRIGHT (c) 2004 The Thomson Corporation.
     STN
                                                            DUPLICATE 6
ΑN
     2003:86974 BIOSIS
DN
     PREV200300086974
TI
     Complete Freund's
                           ***adiuvant***
                                             immunization prolongs survival in
     experimental prion disease in mice.
ΑU
     Tal, Yuval; Souan, Lina; Cohen, Irun R. [Reprint Author]; Meiner, Zeev;
     Taraboulos, Albert; Mor, Felix
     Department of Immunology, Weizmann Institute of Science, P.O. Box 26,
CS
     Rehovot, 76100, Israel
     irun.cohen@weizmann.ac.il
     Journal of Neuroscience Research, (January 15 2003) Vol. 71, No. 2, pp.
SO
     286-290. print.
     ISSN: 0360-4012 (ISSN print).
DT
     Article
```

```
English
ED
    Entered STN: 6 Feb 2003
    Last Updated on STN: 6 Feb 2003
     ANSWER 40 OF 76
                           MEDLINE on STN
     2003475318
                     MEDLINE
٩N
    PubMed ID: 14550926
NC
    Immunisation with a synthetic prion protein-derived peptide prolongs
ΤI
     survival times of mice orally exposed to the scrapic agent.
٩U
    Schwarz Anja; Kratke Oliver; Burwinkel Michael; Riemer Constanze; Schultz
     Julia; Henklein Peter; Bamme Theresa; Baier Michael
CS
    Project Neurodegenerative Diseases, Robert-Koch-Institute, Nordufer 20.
    13353 Berlin, Germany.
Neuroscience letters, (2003 Oct 30) 350 (3) 187-9.
Journal code: 7600130. ISSN: 0304-3940.
50
     Ireland
TC
    Journal; Article; (JOURNAL ARTICLE)
_A
=S
    English
    Priority Journals
ΞΜ
    200311
ΞD
    Entered STN: 20031011
    Last Updated on STN: 20031111
    Entered Medline: 20031110
    ANSWER 41 OF 76 USPATFULL on STN
_3
                                                              DUPLICATE 7
٩N
       2002:272456 USPATFULL
ΓΙ
       Antibodies specific for native
                                           ***PrPSc***
       Prusiner, Stanley B., San Francisco, CA, UNITED STATES
EN
       Williamson, R. Anthony, San Diego, CA, UNITED STATES
       Burton, Dennis R., La Jolla, CA, UNITED STATES
PΙ
                                  20021017
       US 2002150571
                            Α1
       US 6562341
                            В2
                                  20030513
ľ
       US 2001-943906
                                 20010830 (9)
                            Α1
       Continuation of Ser. No. US 2000-550374, filed on 13 Apr 2000, PENDING
RLI
      Continuation of Ser. No. US 1998-36579, filed on 6 Mar 1998, PATENTED Division of Ser. No. US 1996-713939, filed on 13 Sep 1996, PATENTED Continuation-in-part of Ser. No. US 1995-528104, filed on 14 Sep 1995,
       ABANDONED
TC
       Utility
       APPLICATION
_N.CNT 2374
       INCLM: 424/130.100
[NCL
NCL
      NCLM:
              424/130.100
      NCLS:
              424/009.100; 424/009.200; 424/147.100; 435/007.100; 435/070.100;
               435/071.100; 436/503.000; 436/518.000; 436/547.000; 530/387.100
C
       [7]
       ICM: A61K039-395
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
    ANSWER 42 OF 76 USPATFULL on STN
                                                              DUPLICATE 8
١N
      2002:259529 USPATFULL
      Discordant helix stabilization for prevention of amyloid formation
Π
IN
      Johansson, Jan, Stockholm, SWEDEN
ŀΙ
      US 2002143105
                                 20021003
                            Α1
      US 6716589
                            В2
                                 20040406
lΙ
      US 2001-988842
                           Α1
                                 20011119 (9)
      US 2000-253695P
                             20001120 (60)
PRAI
      US 2000-251662P
                             20001206 (60)
      Utility
T
S
      APPLICATION
N.CNT 1541
INCL
      INCLM: 525/054.100
ICL
      NCLM: 435/007.200
C
       [7]
       ICM: C08H001-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
    ANSWER 43 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
    2002:185399 CAPLUS
١N
    136:229029
    Method for precipitating mono and multiple layers of organophosphoric and
Ί
    organophosphonic acids and the salts thereof in addition to use thereof
Ν
    Hofer, Rolf; Pawlak, Michael; Textor, Marcus; Schuermann-Mader, Eveline;
    Ehrat, Markus; Tosatti, Samuele
Α
    Zeptosens A.-G., Switz.
    PCT Int. Appl., 88 pp.
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LA
     German
FAN.CNT 1
     PATENT NO.
                           KIND
                                   DATE
                                                APPLICATION NO.
                                                                          DATE
PΙ
     wo 2002020873
                            Α2
                                   20020314
                                                WO 2001-EP10077
                                                                          20010831
              AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH, CN,
              CO, CR, CU, CZ, DE, DK, DM, DZ, EE, ES, FI, GB, GD, GE, GH, GM,
              HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS,
              LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NO, NZ, PL, PT, RO,
              RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, TZ, UA, UG, US, UZ,
              VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
             GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG .089859 A5 20020322 AU 2001-89859 20010831
     AU 2001089859
     EP 1315968
                                                EP 2001-969680
                                   20030604
                            Α2
                                                                          20010831
          R:
              AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
              IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     US 2003186914
                            Α1
                                   20031002
                                                us 2003-363555
                                                                          20030305
PRAI CH 2000-1732
                            Α
                                   20000905
     WO 2001-EP10077
                            W
                                   20010831
os
     MARPAT 136:229029
L3
     ANSWER 44 OF 76 USPATFULL ON STN
       2002:329478
                     USPATFULL
AΝ
       Novel method for down-regulation of amyloid
TI
       Jensen, Martin Roland, Holte, DENMARK
IN
       Rasmussen, Peter Birk, Frederiksberg, DENMARK
       Nielsen, Klaus Gregorius, Soborg, DENMARK
PΙ
       US 2002187157
                                 20021212
                            Α1
       us 2001-785215
AΤ
                            Α1
                                  20010220 (9)
                             20000221
PRAI
       PA 2000-200000265
       US 2000-186295P
                             20000301 (60)
DT
       Utility
       APPLICATION
FS
LN.CNT 3272
       INCLM: 424/185.100
INCL
       INCLS: 424/085.100; 424/085.200
NCL
               424/185.100
       NCLS:
               424/085.100; 424/085.200
IC
       [7]
       ICM: A61K039-00
       ICS: A61K038-19; A61K038-20
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 45 OF 76 USPATFULL on STN
       2002:251957 USPATFULL
ΑN
ΤI
       Conformational and topological protein regulation
IN
       Lingappa, Vishwanath R., San Francisco, CA, UNITED STATES
       Rutkowski, D. Thomas, San Francisco, CO, UNITED STATES
       Hegde, Ramanugan S., Rockville, MD, UNITED STATES
       US 2002137915
PΙ
                            Α1
                                  20020926
       US 2000-739179
ΑI
                                  20001215 (9)
                            Α1
       US 1999-171012P
                             19991215 (60)
PRAI
       US 1999-172350P
                             19991216 (60)
       Utility
DT
FS
       APPLICATION
LN.CNT
       3036
INCL
       INCLM: 536/023.500
       INCLS: 435/007.210; 435/006.000; 435/325.000; 435/070.210; 435/326.000
NCL
       NCLM:
               536/023.500
              435/007.210; 435/006.000; 435/325.000; 435/070.210; 435/326.000
       NCLS:
IC
       ICM: C12Q001-68
       ICS: G01N033-567; C07H021-04; C12P021-04; C12N005-06
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 46 OF 76 USPATFULL ON STN
L3
ΑN
       2002:178549 USPATFULL
TI
       Vaccine for the prevention and treatment of alzheimer's and amyloid
       related diseases
       Chalifour, Robert, Ile Bizard, CANADA
ΙN
       Hebert, Lise, Brossard, CANADA
       Kong, Xianqi, Dollard-des-Oremaux, CANADA
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CODEN: PIXXD2

Patent

DT

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Gervais, Francine, Ile Bizard, CANADA
       us 2002094335
PΙ
                           Α1
                                 20020718
       us 2001-867847
                           Α1
                                 20010529 (9)
ΑI
RLI
       Continuation-in-part of Ser. No. US 2000-724842, filed on 28 Nov 2000,
       PENDING
       US 1999-168594P
Utility
PRAI
                            19991129 (60)
DT
       APPLICATION
FS
LN.CNT 1946
INCL
       INCLM: 424/185.100
NCL
       NCLM: 424/185.100
       [7]
IC
       ICM: A61K039-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 47 OF 76 USPATFULL ON STN
       2002:99410
                   USPATFULL
ΑN
       Methods and compostions for the treatment and/or diagnosis of
ΤI
       neurological diseases and disorders
IN
       Solomon, Beka, Herzlia Pituach, ISRAEL
       Frenkel, Dan, Rehovot, ISRAEL
       US 2002052311
PΙ
                           Α1
                                 20020502
       us 2001-808037
                           Α1
                                 20010315 (9)
ΑI
       Continuation-in-part of Ser. No. US 2000-629971, filed on 31 Jul 2000,
RLI
       PENDING Continuation-in-part of Ser. No. US 1999-473653, filed on 29 Dec
       1999, PENDING
       US 1999-152417P
Utility
                            19990903 (60)
PRAI
DT
       APPLICATION
FS
LN.CNT 4074
       INCLM: 514/002.000
INCL
       INCLS: 424/093.210
              514/002.000
NCL
       NCLM:
       NCLS: 424/093.210
IC
       [7]
       ICM: A61K048-00
       ICS: A61K038-00
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 48 OF 76 USPATFULL ON STN
       2002:72861 USPATFULL
ΑN
ΤI
       Pharmaceutical compositions comprising a soluble laminin receptor
       precursor or a compound which blocks the interaction of the laminin
                                  ***PrPSc***
       receptor precursor and
                                                 or PrPc
       Weiss, Stefan, Munich, GERMANY, FEDERAL REPUBLIC OF
IN
       US 2002040001
PI
                           Α1
                                 20020404
ΑI
       us 2001-964566
                                 20010928 (9)
                           Α1
       Continuation-in-part of Ser. No. US 2000-424754, filed on 13 Apr 2000,
RLI
       PENDING
PRAI
       EP 1997-108712
                            19970530
       WO 1998-EP3220
                            19980529
DT
       Utility
FS
       APPLICATION
LN.CNT 2050
       INCLM: 514/012.000
INCL
       INCLS: 435/007.100
NCL
              514/012.000
       NCLM:
              435/007.100
       NCLS:
       [7]
IC
       ICM: A61K038-17
       ICS: G01N033-53
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 49 OF 76 USPATFULL ON STN
AN
       2002:297296 USPATFULL
       Methods for inhibition of membrane fusion-associated events, including
TT
       respiratory syncytial virus transmission
       Bolognesi, Dani Paul, Durham, NC, United States
Matthews, Thomas James, Durham, NC, United States
ΙN
       Wild, Carl T., Durham, NC, United States
       Barney, Shawn O'Lin, Cary, NC, United States
       Lambert, Dennis Michael, Cary, NC, United States
       Petteway, Stephen Robert, Cary, NC, United States
       Langlois, Alphonse J., Durham, NC, United States
       Trimeris,
                 Inc., Durham, NC, United States (U.S. corporation)
PA
PΙ
       us 6479055
                           в1
                                20021112
```

```
ΑI
        US 1995-470896
                                    19950606 (8)
        Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994,
RLI
        now patented, Pat. No. US 6017536 Continuation-in-part of Ser. No. US
        1994-255208, filed on 7 Jun 1994 Continuation-in-part of Ser. No. US
        1993-73028, filed on 7 Jun 1993, now patented, Pat. No. US 5464933
        Utility
DT
FS
        GRANTED
LN.CNT 26553
        INCLM: 424/211.100
INCL
        INCLS: 424/186.100; 530/324.000
        NCLM: 424/211.100
NCL
        NCLS: 424/186.100; 530/324.000
IC
        [7]
        ICM: A61K039-145
        435/5; 435/240.2; 424/184.1-189.1; 424/204.1-211.1; 424/225.1; 424/227.1; 424/230.1; 514/1; 514/2; 530/324; 530/350; 530/826
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
      ANSWER 50 OF 76 USPATFULL ON STN
        2002:81024 USPATFULL
ΑN
TI
        Antibodies specific for native
                                              ***PrPSc***
        Prusiner, Stanley B., San Francisco, CA, United States
IN
        Williamson, R. Anthony, San Diego, CA, United States
        Burton, Dennis R., La Jolla, CA, United States
The Regents of the University of California, Oakland, CA, United States
PA
        (U.S. corporation)
        The Scripps Research Institute, La Jolla, CA, United States (U.S.
        corporation)
PΙ
        US 6372214
                                    20020416
                              В1
        us 2000-550374
ΑI
                                    20000413 (9)
        Continuation of Ser. No. US 1998-36579, filed on 6 Mar 1998 Division of
RLI
        Ser. No. US 1996-713939, filed on 13 Sep 1996, now patented, Pat. No. US
                  issued on 8 Dec 1998 Continuation-in-part of Ser. No. US
        1995-528104, filed on 14 Sep 1995, now abandoned
DT
        Utility
        GRANTED
FS
LN.CNT 2518
INCL
        INCLM: 424/130.100
        INCLS: 424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100;
                436/503.000; 436/518.000; 436/547.000; 530/387.100
NCL
                424/130.100
                424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100; 436/503.000; 436/518.000; 436/547.000; 530/387.100
        NCLS:
IC
        [7]
        ICM: A61K039-395
        ICS: C12P019-00; G01N033-567; G01N033-543; C07K016-00
        424/9.1; 424/130.1; 424/147.1; 435/7.1; 435/70.1; 435/71.1; 530/387.1; 436/518; 436/503; 436/547
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 51 OF 76 USPATFULL ON STN
L3
ΑN
        2002:51103
                     USPATFULL
TI
        Methods and compositions for the manufacture of halogenated
        anthracyclines with increased antitumor activity, other anthracyclines,
        halogenated sugars, and glycosyl donors
        Priebe, Waldemar, 4239 Emory St., Houston, TX, United States 77005 Krawczyk, Marta, 175 N. Locust Hill Dr. apt. #2308, Lexington, KY,
IN
        United States
                        40503
        Skibicki, Piotr, Waszyngton Street 39 Apartment 24, Warsaw 04015, POLAND Fokt, Izabela, 1908 Nursery Rd., The Woodlands, TX, United States 77380
        Dziewiszek, Krzysztof, 1908 Nursery Rd., The Woodlands, TX, United
        Grynkiewicz, Grzegorz, .mu.. Zielona 16B/2, 05-092 Lomianki, POLAND
        Perez-Soler, Roman, 564 1st Ave. #20T, New York, NY, United States
        10016
PΙ
        us 6355784
                             в1
                                   20020312
        US 1999-330226
ΑI
                                   19990610 (9)
        US 1998-89162P
PRAI
                              19980612 (60)
        Utility
DT
        GRANTED
LN.CNT 2062
INCL
        INCLM: 536/006.400
        INCLS: 536/004.100; 536/017.200; 536/018.400; 536/018.700; 536/122.000
NCL
        NCLM:
                536/006.400
        NCLS:
                536/004.100; 536/017.200; 536/018.400; 536/018.700; 536/122.000
IC
```

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ICM: C07H015-24
        536/6.4; 536/18.6; 536/122; 536/18.4; 536/18.7; 536/4.1; 536/17.2;
EXF
        514/34
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 52 OF 76 SCISEARCH COPYRIGHT (c) 2004 The Thomson Corporation.
     2002:986508 SCISEARCH
AN
     The Genuine Article (R) Number: 620PU
GA
TI
     Induction of antibodies against murine full-length prion protein in
     wild-type mice
     Koller M F; Grau T; Christen P (Reprint)
ΑU
     Univ Zurich, Inst Biochem, Winterhurerstr 190, CH-8057 Zurich, Switzerland
CS
     (Reprint); Univ Zurich, Inst Biochem, CH-8057 Zurich, Switzerland
CYA
     Switzerland
     JOURNAL OF NEUROIMMUNOLOGY, (NOV 2002) Vol. 132, No. 1-2, pp. 113-116. Publisher: ELSEVIER SCIENCE BV, PO BOX 211, 1000 AE AMSTERDAM,
     NETHERLANDS.
     ISSN: 0165-5728.
     Article; Journal
DT
     English
LA
REC
     Reference Count: 17
     *ABSTRACT IS AVAILABLE IN THE ALL AND IALL FORMATS*
L3
     ANSWER 53 OF 76 USPATFULL ON STN
                                                            DUPLICATE 9
ΑN
       2001:224147 USPATFULL
        Prevention of cancer
TI
IN
       Henrik, Raskov Hans, Hellerup, Denmark
       us 2001049364
                                  20011206
PΙ
                           Α1
       US 6703380
                            В2
                                  20040309
       US 2001-825891
ΑT
                          Α1
                                 20010405 (9)
       Continuation-in-part of Ser. No. WO 2000-DK546, filed on 29 Sep 2000,
RLI
       UNKNOWN
PRAI
       DK 1999-1390
                             19990929
DT
       Utility
       APPLICATION
FS
LN.CNT
       1937
       INCLM: 514/164.000
INCL
       INCLS: 514/167.000
NCL
               514/165.000
       NCLM:
               514/167.000
       NCLS:
IC
        [7]
       ICM: A61K031-616
       ICS: A61K031-59
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
     ANSWER 54 OF 76 CAPLUS COPYRIGHT 2004 ACS ON STN
L3
ΑN
     2001:816487
                  CAPLUS
DN
     135:356752
     Epitope synchronization in antigen presenting cells
TT
IN
     Simard, John J. L.; Diamond, David C.; Lei, Xiang-Dong
     CTL Immunotherapies Corp., USA
PA
SO
     PCT Int. Appl., 131 pp.
     CODEN: PIXXD2
DT
     Patent
     English
LA
FAN.CNT 7
     PATENT NO.
                           KIND
                                  DATE
                                               APPLICATION NO.
                                                                        DATE
                           ____
PΙ
     wo 2001082963
                           A2
                                  20011108
                                               WO 2001-US13806
                                                                        20010427
     wo 2001082963
                           Α3
                                  20020411
             AE, AG, AL, AM, AT, AT, AU, AZ, BA, BB, BG, BR, BY, BZ, CA, CH,
              CN, CR, CU, CZ, CZ, DE, DE, DK, DK, DM, DZ, EE, EE, ES, FI, FI,
              GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, JP, KE, KG, KP, KR,
              KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ,
              NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SK, SL, TJ, TM, TR,
              TT, TZ,
                      UA, UG, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ, MD, RU,
              TJ.
         RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF,
              BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG
                                  20030122
                                               EP 2001-930922
                            Α2
                                                                        20010427
             AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
              IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
     JP 2003535824
                           T2
                                  20031202
                                               JP 2001-579836
                                                                        20010427
PRAI US 2000-560465
                                  20000428
                            Α
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us 2000-561074
                                     20000428
     US 2000-561571
                              Α
                                     20000428
     us 2000-561572
                              Α
                                     20000428
     wo 2001-US13806
                                     20010427
     ANSWER 55 OF 76 CAPLUS COPYRIGHT 2004 ACS ON STN
L3
     2001:507867 CAPLUS
ΑN
     135:91527
DN
     Tissue-specific DNA delivery via M cell-directed vaccines, and enhanced in
TI
     vivo mucosal IgA and T cell responses resulting therefrom
IN
     Pascual, David W.
     Research and Development Institute, Inc., USA
PA
SO
     PCT Int. Appl., 58 pp.
     CODEN: PIXXD2
DT
     Patent
LA
     English
FAN.CNT 3
     PATENT NO.
                                                  APPLICATION NO.
                            KIND
                                     DATE
                                                                             DATE
     wo 2001049867
                                     20010712
                                                  wo 2001-US426
                             Α1
                                                                             20010108
               AE, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BY, CA, CH, CN, CU, CZ,
               DE, DK, EE, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS,
               JP, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MD, MG, MK,
               MN, MW, MX, NO, NZ, PL, PT, RO, RU, SD, SE, SG, SI, SK, SL, TJ, TM, TR, TT, UA, UG, US, UZ, VN, YU, ZA, ZW, AM, AZ, BY, KG, KZ,
                  RU,
                        TJ, TM
          RW: GH, GM, KE, LS, MW, MZ, SD, SL, SZ, TZ, UG, ZW, AT, BE, CH, CY, DE, DK, ES, FI, FR, GB, GR, IE, IT, LU, MC, NL, PT, SE, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GW, ML, MR, NE, SN, TD, TG L257654 A1 20021120 EP 2001-901811 20010108
     EP 1257654
              AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT,
               IE, SI, LT, LV, FI, RO, MK, CY, AL, TR
                                                  us 2002-169492
     us 2004033486
                              Α1
                                     20040219
                                                                             20021021
     US 2004109871
                              Α1
                                                  us 2003-660787
                                                                             20030912
                                     20040610
PRAI US 2000-174786P
                             Ρ
                                     20000106
     WO 2001-US426
                              W
                                     20010108
         2001-274639P
     US
                              Ρ
                                     20010312
     WO 2002-US7254
                             A2
                                     20020312
     US 2002-169492
                                     20021021
                             A2
RE.CNT
                THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD
                ALL CITATIONS AVAILABLE IN THE RE FORMAT
L3
     ANSWER 56 OF 76 USPATFULL ON STN
        2001:157792 USPATFULL
ΑN
                                              ***PrPSc***
TI
        Antibodies specific for native
IN
        Prusiner, Stanley B., San Francisco, CA, United States
        Williamson, R. Anthony, San Diego, CA, United States
        Burton, Dennis R., La Jolla, CA, Únited States
The Scripps Research Institute, La Jolla, CA, United States (U.S.
PA
        corporation)
PΙ
        us 6290954
                              в1
                                    20010918
AΙ
        US 1998-36579
                                    19980306 (9)
RLI
        Division of Ser. No. US 1996-713939, filed on 13 Sep 1996, now patented,
        Pat. No. US 5846533 Continuation-in-part of Ser. No. US 1995-528104,
        filed on 14 Sep 1995, now abandoned
DT
        Utility
FS
        GRANTED
LN.CNT
       2513
        INCLM: 424/130.100
INCL
        INCLS: 424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100;
                436/503.000; 436/518.000; 436/547.000; 530/387.100
                424/130.100
NCL
        NCLS:
                424/009.100; 424/147.100; 435/007.100; 435/070.100; 435/071.100;
                436/503.000; 436/518.000; 436/547.000; 530/387.100
IC
        [7]
        ICM: A61K039-395
        ICS: G01N033-53; G01N033-567; C07K016-00
        424/9.1; 424/130.1; 424/147.1; 435/7.1; 435/70.1; 435/71.1; 530/387.1;
EXF
        436/518; 436/503; 436/547
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
     ANSWER 57 OF 76 USPATFULL on STN
        2001:125727 USPATFULL
AN
TI
        Correction of genetic defects using chemical chaperones
        Welch, William J., San Francisco, CA, United States
Brown, C. Randell, Hershey, PA, United States
ΙN
```

```
Tatzelt, Jorg, Munchen, Germany, Federal Republic of
        The Regents of The University of California, Oakland, CA, United States
PA
        (U.S. corporation)
        us 6270954
PΙ
                               В1
                                     20010807
        US 1999-291406
ΑI
                                     19990413 (9)
        Continuation-in-part of Ser. No. US 1997-838691, filed on 9 Apr 1997,
RLI
        now patented, Pat. No. US 5900360
        US 1996-15155P
                                19960410 (60)
PRAI
        Utility
DT
FS
        GRANTED
LN.CNT 2342
        INCLM: 435/004.000
INCL
        INCLS: 435/026.000; 435/023.000; 435/024.000; 435/963.000
NCL
                435/004.000
        NCLM:
                435/023.000; 435/024.000; 435/026.000; 435/963.000
        NCLS:
IC
        [7]
        ICM: C12Q001-00
        ICS: C12Q001-32; C12Q001-37
        435/4; 435/26; 435/23; 435/24; 435/963
FXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 58 OF 76 USPATFULL on STN
L3
        2001:67794 USPATFULL
ΑN
        Human respiratory syncytial virus peptides with antifusogenic and
TI
        antiviral activities
        Barney, Shawn O'Lin, Cary, NC, United States
Lambert, Dennis Michael, Cary, NC, United States
Petteway, Stephen Robert, Cary, NC, United States
Trimeris, Inc., Durham, NC, United States (U.S. corporation)
IN
PA
        US 6228983
                                     20010508
PΙ
                               В1
        us 1995-485264
                                     19950607 (8)
ΑI
RLI
        Division of Ser. No. US 1995-470896, filed on 6 Jun 1995
        Continuation-in-part of Ser. No. US 1994-360107, filed on 20 Dec 1994
        Continuation-in-part of Ser. No. US 1994-255208, filed on 7 Jun 1994 Continuation-in-part of Ser. No. US 1993-73028, filed on 7 Jun 1993, now
        patented, Pat. No. US 5464933
DT
        Utility
        Granted
FS
LN.CNT 32166
INCL
        INCLM: 530/300.000
        INCLS: 530/324.000; 530/325.000; 530/326.000; 424/211.100; 424/186.100
NCL
                 530/300.000
        NCLS:
                424/186.100; 424/211.100; 530/324.000; 530/325.000; 530/326.000
        [7]
IC
        ICM: A61K038-00
EXF
        530/350; 530/324-329; 530/300; 424/211.1
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
L3
      ANSWER 59 OF 76 USPATFULL ON STN
ΑN
        2001:51590 USPATFULL
        Clearance and inhibition of conformationally altered proteins
TI
IN
        Prusiner, Stanley B., San Francisco, CA, United States
        Supattapone, Surachai, San Francisco, CA, United States
        Scott, Michael, San Francisco, CA, United States
PA
        The Regents of the University of California, Oakland, CA, United States
        (U.S. corporation)
PΙ
        US 6214366
                                     20010410
                               В1
        US 1999-322903
ΑI
                                     19990601 (9)
DT
        Utility
FS
        Granted
LN.CNT 1037
INCL
        INCLM: 424/405.000
        INCLS: 424/438.000; 424/442.000; 424/484.000; 424/DIG.016; 424/078.320;
                424/078.350; 424/078.360; 424/078.370; 424/078.380; 514/772.300; 514/772.400; 514/772.500; 514/772.600; 514/772.700
                424/405.000
NCL
        NCLM:
                424/078.320; 424/078.350; 424/078.360; 424/078.370; 424/078.380; 424/438.000; 424/442.000; 424/484.000; 424/DIG.016; 514/772.300; 514/772.400; 514/772.500; 514/772.600; 514/772.700
        NCLS:
IC
        [7]
        ICM: A01N025-10
        424/78.32; 424/78.35; 424/78.38; 424/405; 424/438; 424/442; 424/DIG.16; 514/772.3-772.7
EXF
CAS INDEXING IS AVAILABLE FOR THIS PATENT.
      ANSWER 60 OF 76 WPIDS COPYRIGHT 2004 THOMSON DERWENT ON STN
L3
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```
2002-049350 [06]
ΑN
                         WPIDS
     N2002-036483
                         DNC C2002-013896
DNN
     New polypeptides,
                       useful as antiviral agents, comprise their prion
TT
     proteins able to bind nucleic acid, nucleocapsid proteins, and ligands for
     use as antiprion agents.
     B04 C06 D16 S03
DC
     DARLIX, J L; GABUS, D C; LEBLANC, P; DARLIX, J; GABUS-DARLIX, C
ΙN
     (INRM) INSERM INST NAT SANTE & RECH MEDICALE
PΑ
CYC
PΙ
     wo 2001083747
                      A2 20011108 (200206)* FR
                                                         C12N015-12
        RW: AT BE CH CY DE DK EA ES FI FR GB GH GM GR IE IT KE LS LU MC MW MZ
            NL OA PT SD SE SL SZ TR TZ UG ZW
         W: AE AG AL AM AT AU AZ BA BB BG BR BY BZ CA CH CN CO CR CU CZ DE DK
            DM DZ EE ES FI GB GD GE GH GM HR HU ID IL IN IS JP KE KG KP KR KZ
             LC LK LR LS LT LU LV MA MD MG MK MN MW MX MZ NO NZ PL PT RO RU SD
             SE SG SI SK SL TJ TM TR TT TZ UA UG US UZ VN YU ZA ZW
                      A1 20011102 (200206)
     FR 2808278
                                                         C07K014-435
     AU 2001056454
                        20011112 (200222)
                                                         C12N015-12
ADT
     WO 2001083747 A2 WO 2001-FR1336 20010430; FR 2808278 A1 FR 2000-5535
     20000428; AU 2001056454 A AU 2001-56454 20010430
FDT
     AU 2001056454 A Based on WO 2001083747
                           20000428
PRAI FR 2000-5535
          C07K014-435; C12N015-12
IC
     ICM
          A61K038-17; A61K039-395; A61K048-00; A61P025-00; A61P031-12; C07K014-47; C07K019-00; C12N007-00; C12N015-86; C12Q001-68; C12Q001-70; G01N033-569; G01N033-68
     ICS
13
     ANSWER 61 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.
     on STN
                                                           DUPLICATE 10
     2001435095
                 EMBASE
AN
     Human aqueous humor levels of oral ciprofloxacin, levofloxacin, and
ΤI
     moxifloxacin.
ΔU
     Garcia-Saenz M.C.; Arias-Puente A.; Fresnadillo-Martinez M.J.;
     Carrasco-Font C.
     Dr. M.C. Garcia-Saenz, Fundacion Hospital de Alcorcon, Budapest 1, 28922
     Alcorcon, Madrid, Spain. mcgarcias@fhalcorcon.es
SO
     Journal of Cataract and Refractive Surgery, (2001) 27/12 (1969-1974).
     Refs: 23
     ISSN: 0886-3350
                      CODEN: JCSUEV
PUI
     s 0886-3350(01)00997-x
CY
     United States
     Journal; Article
DT
     012
FS
             Ophthalmology
     037
             Drug Literature Index
     English
     English
L3
     ANSWER 62 OF 76 EMBASE COPYRIGHT 2004 ELSEVIER INC. ALL RIGHTS RESERVED.
     on STN
AN
     2003330992 EMBASE
TI
     Short-term effect of mitomycin-C augmented trabeculectomy on axial length
     and corneal astigmatism.
ΑU
     Kook M.S.; Kim H.B.; Lee S.U.
     Dr. M.S. Kook, Department of Ophthalmology, Ulsan University School of
CS
     Medicine, Asan Medical Center, 388-1 Pungnap-dong, Songpa-gu, Seoul,
     138-736, Korea, Republic of
SO
     Journal of Cataract and Refractive Surgery, (2001) 27/4 (518-523).
     Refs: 25
     ISSN: 0886-3350
                      CODEN: JCSUEV
PUI
     s 0886-3350(00)00646-5
CY
     United States
DT
     Journal; Article
FS
     012
             Ophthalmology
     037
             Drug Literature Index
     038
             Adverse Reactions Titles
     English
LA
SI
     English
L3
     ANSWER 63 OF 76 CAPLUS COPYRIGHT 2004 ACS on STN
     2000:861780
AN
                  CAPLUS
DN
     134:28436
TI
     Vaccines against conformation-dependent protein and non-protein antigens
TN
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PA
     Max-Delbruck-Centrum fuer Molekulare Medizin, Germany
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     Ministry of Agriculture, Foresty and Fisheries National Institute of
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     Pharmaceutical composition comprising immunogen against amyloid component such as fibril peptide or protein, or antibody against amyloid component
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       Finlay J.L.; Abramson D.H.
       Dr. I.J. Dunkel, Department of Pediatrics, Mem. Sloan-Kettering Cancer
CS
       Center, Box 185, 1275 York Avenue, New York, NY 10021, United States.
       E-mail: dunkeli@mskcc.org
S0
       Cancer, (15 \text{ NOV } 2000), 89/10 (2117-2121), 17 \text{ reference}(s)
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Brown, C. Randell, 1470 9th Ave. #12, San Francisco, CA, United States
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        Tatzelt, Jorg, 740 Parnassus, San Francisco, CA, United States 94122
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       for nephroblastoma initially treated according to the SIOP 9/GPOH trial
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       Dept. of Pediatric Hematology and Oncology, University of Jena, Germany,
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       Federal Republic of
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     Journal of Clinical Oncology, (1989) vol. 7, No. 12, pp. 1824-1830.
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      with High Dose Chemotherapy with Autologous Stem Cell Rescue (
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      Williams S; Bitran J; Desser R; Golick J; Beschorner J; Fullem L
      Chicago, Illinois, United States
      Proc.Am.Soc.Clin.Oncol. (7, 24 Meet., 9, 1988)
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